Acute effect of primary percutaneous coronary intervention on left ventricular dyssynchrony in ST-segment elevation myocardial infarction

Sinan İnci, Şule Karakelleoğlu, M. Hakan Taş, Şakir Arslan, Fuat Gündoğdu, Eftal Murat Bakırcı, Hüsnü Değirmenci, Hüseyin Senocak

Department of Cardiology, Faculty of Medicine, Atatürk University; Erzurum-Turkey

ABSTRACT

Objective: The aim of this study was to prospectively evaluate the effect of percutaneous coronary intervention in the acute period on left ventricular dyssynchrony in ST-segment elevation myocardial infarction patients by using Tissue Synchronization Imaging.

Methods: Forty-four ST-segment elevation myocardial infarction (MI) patients (29 male, 15 female), who were admitted within the first 12 hours of chest pain symptoms, were enrolled in the study. According to the localization of MI, the patients were divided into groups as anterior MI (n=26) and inferior MI (n=18). All echocardiography measurements were taken just before percutaneous coronary intervention (PCI) and following PCI at a mean of 3-6 days. They were assessed according to the time to reach the peak systolic velocity, which was calculated by the tissue synchronization imaging method from four pairs of non-apical alternate segments. The difference between the duration to reach the peak systolic velocity in alternate segments was regarded as left ventricle dyssynchrony and the results were compared.

Results: In the anterior MI group, basal anterior (p<0.01), mid-anterior segment (p<0.01) and basal septal segment (p<0.01); in the inferior MI group, the basal septal segment (p=0.02), mid-septal segment (p=0.02), and basal and mid-inferior segment (p<0.01) values were significantly lower in the post-PCI measurements when compared to the measurements taken prior to PCI. In both groups, the intraventricular dyssynchrony indices of the basal anterior-basal inferior (p<0.01), mid-anterior-mid-inferior (p<0.01) segments were found to be significantly lower in the post- PCI period when compared to the pre-PCI period.

Conclusion: It was found that percutaneous coronary intervention in patients with ST-elevation significantly decreases the degree of LV dys-synchrony in the acute period. (Anadolu Kardiyol Derg 2014; 14: 591-8)

Key words: percutaneous coronary intervention, dyssynchrony, tissue synchronization imaging, ST-segment elevation myocardial infarction

Introduction

Mechanical synchronization disorder leads to a decrease in ejection fraction and stroke volume, an abnormal distribution in wall tension, and more workload during cardiac contraction (1). Various techniques to assess dyssynchronization have been developed (2-4). Tissue Doppler echocardiography is one of the auxiliary echocardiography techniques that is performed to evaluate mechanical dyssynchronization. Tissue Synchronization Imaging (Tissue Synchronization Imaging TM, GE Vingmed, Horten, Norway) is a tissue Doppler echocardiography program developed in 2004 (5).

Delayed electrical activation of one of the myocardial walls results in a mechanical dyssynchrony. In a normal left ventricle,

all segments are activated and thus contract at the same time. When the septum begins to contract, it contributes to an increase in left ventricle (LV) pressure and ejection. The combination of early septal contraction stretching in the lateral wall and late lateral wall contraction stretching of the septum results in a reduced systolic function, and causes a dissipation of contractile forces in the LV (6).

The relationship between heart failure and myocardial synchronization have been documented with much pathologic and physiologic evidence (7-11). When the heart is functioning normally, there must be a synchronized movement between the atrium and the ventricle, the right and left ventricles, and various segments of the ventricles for blood to be pumped normally. This normal synchronization is dependent on a normal conduction



system, stimulation-contraction, myocardial structure, and heart function (12). A similar relation between heart failure and myocardial synchronization has been shown in cardiac diseases such as acute myocardial infarction (2-4, 13), unstable angina pectoris (14, 15) and hypertrophic cardiomyopathy (16). The studies that demonstrate the effect of percutaneous coronary intervention on myocardial synchrony in acute myocardial infarction are limited in number. The aim of the current study was to show the effect of percutaneous coronary intervention (PCI) in the acute period on left ventricular dyssynchrony in ST-segment elevation myocardial infarction patients using Tissue Synchronization Imaging (TSI).

Methods

Study population

This study was performed in our clinics between April 2008 and June 2010. Forty-four patients (29 male, 15 female), who were diagnosed with acute ST-segment elevation acute myocardial infarction (AMI) according to ESC/ACC criteria (17), presented to our clinic within the first twelve hours of the onset

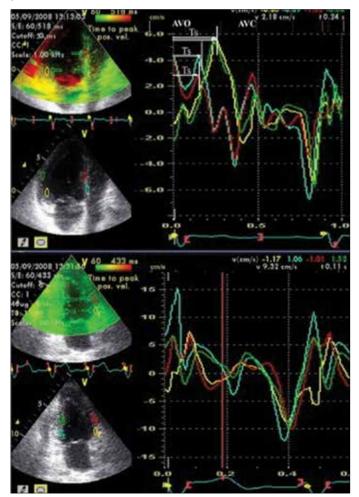


Figure 1. Graphic showing the duration to peak systol and apical four cavity and two cavity TSI images of a patient with acute MI

AVC - aort valve closure; AVO - aort valve opening; TSI-tissue synchronization imaging

of AMI, and who had an indication for PCI were included in this study. Before the measurements were taken according to the infarction area, the patients were divided into two groups as anterior AMI and inferior AMI.

Exclusion criteria:

Exclusion criteria: Previously diagnosed with heart failure, QRS duration exceeding 120 msc, cardiogenic shock, serious valvular heart disease, infarction history, uncontrolled hypertension, hypertrophic obstructive cardiomyopathy.

Patients with multiple vascular disease at coronary angiography;

Atrial fibrilasyon, prosthetic valve, pace rhythm, who had tachycardia during measuments, poor image quality, systemic diseases, chronic kidney disease.

The study was conducted in accordance with the principles of the Declaration of Helsinki and was approved by the local Ethic Committee. All patients and the healthy controls were informed about the study and their written consent forms were obtained.

Echocardiography

The first echocardiographic recordings of all the patients were obtained just before PCI in the catheterization laboratory. Standard transthoracic echocardiography measurements of all patients were taken by using a 2.5 Mhz probe of a Vivid 7 device (GE Healthcare). All images were recorded as three strokes on average for off-line measurements using the protocol of the Echopack program. Left ventricular ejection fractions were measured by a modified Simpson's technique, and left ventricular diastolic diameter, systolic diameter, interventricular septum thickness, and posterior wall thickness were measured by using M-mode.

Tissue synchronization imaging

In the tissue Doppler imaging mode, the cursor was placed in the basal and middle regions of the lateral, septal, anterior, and inferior walls. Myocardial velocity graphics were obtained from apical-four cavity images and apical two cavity images. The apical segments of the heart were not taken into consideration in TSI since they were not coded in color in the early phases the starting point was set at the point of the opening of the aortic valves, and the end point was set at 200 milliseconds after the closure of the aortic valves, the beginning of the rapid filling period. The duration from the start of QRS to peak systolic velocity (Tscore) was measured in 8 segments (Fig. 1). The interobserver and intraobserver variabilities have been compared in 60 consecutive measurements and were 4.7% and 3.2%, respectively. Bazett's formula was applied to remove the effect of heart rate variation on Tscore. Measurements were taken by the Bazett formula according to the heart rate with the following formula (12):

Ts=Tscore/ \sqrt{R} -R (Bazett formula)

The absolute value of the Tscor differences between alternate segments (inferior-anterior and lateral-septal) was regard-

Table 1. Clinical characteristics of the patients (n=44)

Age, years	58.4±9.9
Male gender (%) (n)	66 (29)
Female, gender (%) (n)	34 (15)
Diagnosis	
Anterior MI (%) (n)	59. (26)
Inferior MI (%) (n)	41 (18)
Diabetes mellitus (%) (n)	27 (12)
Cigarette smoking (%) (n)	56 (25)
Family history (%) (n)	16 (7)
Hypertension (%) (n)	68 (29)
Hyperlipidemia (%) (n)	21 (9)
Time of acceptance (hour)(min-max duration)	5.1±1.1 (1-11)
Systolic blood pressure, mm Hg	115.4±16.3
Diastolic blood pressure, mm Hg	73.1±10.1
Heart rate, strokes / min	78.6±15.8
Peak CK (U/I)	1447.9±1323.5
Peak CKMB (U/I)	244.2±302.7
Peak troponin, μg/l	14.9±9.2
Hospitalization, day	4.8±2.4
Medical treatment (%) (n)	
ASA	98 (43)
Clopidogrel	100 (44)
LMWH	91 (40)
Beta blocker	87 (38)
ACE inh	93 (41)
Digoxin	16 (7)
Statin	89 (39)
Diuretics	32 (14)
Type of Lesion that received intervention (%)	(n)
LAD	59 (25)
RCA	34 (15)
CX	7 (3)
Additional lesion (%) (n)	45.5

ACE in - angiotensin converting enzyme inhibitors; ASA - acethylsalicyclic acid; CK - creatinine kinase; Cx - left circumflex coronary artery; LAD - left anterior descending coronary artery; LMWH - low molecular weight heparin; MI - myocardial infarction; RCA - right coronary artery

ed as intraventricular dyssynchrony (18). Intraventricular dyssynchrony before and after PCI were calculated separately. Changes in dyssynchronization values prior to and following PCI (mean of 3-6 days) were compared in each patient to identify the differences.

Statistical analysis

All parameters were expressed as mean± standard deviation. All data were analyzed by using SPSS for Windows version

15.0 software (Chicago, IL, USA). Categorical variables were presented as frequencies and percentages continuous variables were expressed as means and SD. The normal distribution of continuous variables was tested with Kolmogorov-Smirnov test . Wilcoxon test was used to find the statistical differences in the patient groups. P values <0.05 were considered statistically significant.

Results

Clinical characteristics

The clinical and laboratory features of the study population are summarized in Table 1. Twenty-nine of the 44 patients that were enrolled in the study were male (65.9%). The average age of the patients was 58.4±9.9 years. Twenty-six of the patients (59.1%) had a diagnosis of anterior AMI and 18 were diagnosed with inferior AMI. Twelve of the patients (27.3%) had diabetes mellitus (DM), 30 (68.2%) had hypertension (HT), 25 had (56.8%) cigarette smoking history, 9 (20.5%) had hyperlipidemia, and 7 (15.9%) had a family history. Patients were admitted to our clinics within an average of 5.1±1.1 hours after the onset of chest pain. Percutaneous coronary intervention was performed on left anterior descending (LAD) artery lesions in 26 patients (59.1%), right coronary artery (RCA) lesions in 15 patients (34.1%), and left circumflex (CX) lesions in 3 patients (6.8%). Medical treatment that the patients had taken throughout the duration of their hospital stay is shown in Table 1.

Effects of PCI on the conventional echocardiographic parameters

Standard transthoracic echocardiography data of the patients were recorded at the time of referral (pre- PCI) and right before discharge (mean of 3-6 days). Left ventricular systolic diameter (LVSD) and left ventricular diastolic diameter (LVDD), left atrium dimensions, left ventricle ejection fractions (Modified Simpson), interventricular septum thickness (IVS), and posterior wall thickness (PW) were all evaluated. Pre- and post-PCI measurements were found as follows: LVSD [(34.3, 32.8) p<0.01] and LVDD [(48.5, 47.1) p<0.01], respectively. It was documented that post-PCI measurements were significantly lower compared to pre-PCI measurements. The ejection fraction was found to be significantly higher in post-PCI measurements [(37.7%, 42.9) p<0.01]. Other pre and post-PCI measurements were as follows: left atrium dimension [(38.3, 39.3%) p=0.6], interventricular septum thickness [(11.7, 11.8) p=0.5] and posterior wall thickness [(11.7, 11.7) p=0.9]. There were no significant differences between the pre- and post-PCI values of these parameters. Standard transthoracic echocardiography data of the patients are shown in Table 2.

Effects of PCI on the Ts and dyssynchrony indexes

Pre- and post-PCI Ts values were measured in 8 segments. Corrections were made into two groups as the anterior AMI group and inferior AMI group according to the infarction area. In the

Table 2. Standard transthoracic echocardiographic measurements of the patients

	Pre-PCI (n=44)	Post-PCI (n=44)	P
LVDD, mm	48.5	47.1	<0.01
LVSD, mm	34.3	32.8	<0.01
LVEF (%)	37.7	42.9	<0.01
PW, mm	11.7	11.7	NS
IVS, mm	11.8	11.7	NS
LA, mm	38.3	39.3	NS

IVS - interventricular septum thickness; LA - left atrium diameter; LVDD - left ventricular diastolic diameter; LVEF - left ventricular ejection fraction (Modified Simpson method); LVSD - left ventricular systolic diameter; NS - not significant; PW - posterior wall thickness

Table 3. Average Ts values and standard deviations of segments in anterior AMI patients

	Pre-PCI (n=26)	Post-PCI (n=26)	P
Basal-lateral, ms	106.6±15.5	105.8±16.1	0.35
Basal-septal, ms	129.4±36.6	114.1±22.4	<0.01
Mid-lateral, ms	104.9±14.9	104.5±15.1	0.29
Mid-septal, ms	117.5±19.2	113.8±18.8	0.08
Basal-anterior, ms	174.1±18.8	124.7±17.6	<0.01
Basal-inferior, ms	106.3±17.8	105.2±17.1	0.18
Mid-anterior, ms	172.1±19.2	122.3±16.9	<0.01
Mid-inferior, ms	104.5±14.9	104.1±15.0	0.65
Ms - milliseconds; PCI - percutaneous coronary intervention			

Table 4. Pre- and post- PCI intraventricular dyssynchrony values in anterior AMI patients

	Pre-PCI (n=26)	Post-PCI (n=26)	P
MAMI difference, ms	67.5±16.1	19.1±13.3	<0.01
BABI difference, ms	67.7±15.2	19.5±12.5	<0.01
BLBS difference, ms	19.1±14.7	19.2±13.7	0.78
MLMS difference, ms	22.1±14.0	22.1±13.9	0.92

BABI - basal anterior - basal inferior; BLBS - basal lateral - basal septal; MAMI - midanterior - mid inferior; MLMS - mid-lateral - mid-septal; Ms - milliseconds

anterior AMI group, the Ts values of the basal-septal segment, basal-anterior segment, mid-anterior segment were significantly decreased in post-PCI measurements (p<0.01). No significant difference was observed in the Ts values of the basal-lateral segment, mid-lateral segment, mid-septal segment, basal-inferior segment, or the mid-inferior segment in post-PCI measurements (Table 3). In the inferior AMI group, Ts values of the basal-septal segment, mid-septal segment, basal-inferior segment, and mid-inferior segment significantly decreased in post-PCI measurements (p<0.01). No statistical significance was observed in the Ts values of the basal-lateral segment, mid-lateral segment, basal-anterior segment, or the mid-anterior segment in post-PCI measurements (Table 4). Patients were pre- and post-PCI assessed in regard to intraventricular dyssynchronization. The basal-anterior-

Table 5. Average Ts values and standard deviations of segments in inferior AMI patients

	Pre-PCI (n=18)	Post-PCI (n=18)	P
Basal-lateral, ms	106.5±26.8	102.6.4±24.1	0.22
Basal-septal, ms	120.2±34.1	109.7±28.5	0.03
Mid-lateral, ms	105.9±27.4	101.7±21.2	0.22
Mid-septal, ms	113.2±29.7	108.1±29.4	0.04
Basal-anterior, ms	112.8±24.9	112.1±24.1	0.84
Basal-inferior, ms	172.6±43.1	124.4±33.8	<i>P</i> <0.01
Mid-anterior, ms	111.2±25.6	110.1±25.5	0.24
Mid-inferior, ms	169.6±42.8	122.5±33.7	<i>P</i> <0.01
Ms - milliseconds; PCI - percutaneous coronary intervention			

Table 6. Pre- and post- PCI intraventricular dyssynchrony values in inferior AMI patients

	Pre-PCI (n=18)	Post-PCI (n=18)	P
MIMA difference, ms	61.4±28.5	16.6±20.0	<0.01
BIBA difference, ms	60.1±31.3	18.1±20.3	<0.01
BLBS difference, ms	24.5±21.8	22.6±20.4	0.07
MLMS difference, ms	13.7±10.1	12.5±10.1	0.07

BIBA - basal inferior - basal anterior; BLBS - basal lateral - basal septal; MIMA - mid-inferior - mid-anterior; MLMS - mid-lateral - mid-septal; Ms - milliseconds

basal-inferior (BABI) (p<0.01), mid-anterior-mid-inferior (MAMI) (p<0.01) intraventricular dyssynchrony index in both Anterior AMI group and Inferior AMI group were significantly decreased in post-PCI measurements when compared with the pre-PCI measurements (p<0.01). No significant difference was observed in the basal lateral-basal septal (BLBS) and mid-lateral-mid-septal (MLMS) intraventricular dyssynchronization values (Table 5, 6).

Discussion

The current study assessed the effect of percutaneous coronary intervention on acute left ventricular dyssynchrony in ST-segment elevation myocardial infarction. Although the measurements in this study were obtained a short while after percutaneous coronary intervention, there was a significant decrease in LV dyssynchronization after PCI. Although there are many studies in the literature that investigate the effect of PCI on ST-segment elevation myocardial infarction, there aren't studies that assess its effect on left ventricular dyssynchronization.

Echocardiography with tissue Doppler imaging (TDI) is the most commonly used method for the detection of LV dyssynchrony. Arita et al. (19) reported a significant increase in LV dyssynchrony, which was demonstrated by calculating the standard deviation of the time to peak radial strain at six mid-ventricular segments in canine models with heart failure. Karakaş et al. (20) have demonstrated the relation between non-dipper blood pressure and LV dyssynchrony in both normotensive and hypertensive individuals by using tissue Doppler imaging method. In

another study done by Karakas et al. (21), by using the same method they have demonstrated that left ventricular systolic dyssynchrony is an early manifestation of heart involvement in SCA patients with normal EF and narrow QRS. Schiller et al. (22) used the TSI method to evaluate the myocardial dyssynchrony before and after the cardiac pacemaker treatment. They reported that the TSI method is an easy and applicable method in the quantitative detection of regional dyssynchrony. Penicka et al. (23) used the comparison of left ventricle segments' duration to reach peak systolic velocity from the start of QRS to evaluate intraventricular synchronization disorder. It was reported that comparing the durations needed for two alternate walls from four apical cavities to reach peak systolic velocity (septum lateral wall delay) was satisfactory. Thus, it was claimed that a minimum delay of 60 milliseconds is sufficient for the diagnosis of distinct intraventricular mechanical synchronization disorder. Yu et al. (18) evaluated more segments together and expressed the temporal heterogeneity of segmental movement as the standard deviation of 12 basal and middle segments. They determined that having a standard deviation exceeding 33 milliseconds is an important indicator for dyssynchrony. This index, which is called the Yu index, shows a high correlation with the delay between the septum and posterior wall and parasternal short axis. In the current study, the comparison of durations that are passed till the two alternate walls reach peak systolic velocity were performed in four and two apical cavity images (septum-lateral wall delay, anterior- inferior wall delay); peak systolic velocity differences between the walls were used as intraventricular dyssynchrony indexes. In both inferior AMI patients and anterior AMI patients, before percutaneous coronary intervention, in the comparison of anterior and inferior walls, the average intraventricular dyssynchrony index was found to be higher than 60 milliseconds. Following percutaneous coronary intervention, this parameter has an approximate average of 20 milliseconds. As a result, the most distinct finding of this study, viewed in light of the present literature, is that there is significant LV dyssynchrony before percutaneous coronary intervention of acute AMI and that the dyssynchrony degree decreases significantly following percutaneous coronary intervention.

Previous studies suggested the presence and clinical relevance of LV dyssynchrony in the setting of chronic heart failure. In this group of patients, the loss of LV synchronous contraction was related to impaired LV systolic function and poor hemodynamic status, which is an indicator of a poor outcome (11, 24). Cho et al. (25, 26) demonstrated that mechanical dyssynchrony was a powerful predictor of mortality or cardiac events in heart failure patients with normal and wide QRS. Penicka et al. (11), Fauchier et al. (27), and Bader et al. (10) reported that LV dyssynchrony was prognostic of cardiac endpoints. Kutyifa et al. (28) investigated VT/VF events and LV dyssynchrony in mild heart failure patients with LBBB and an implanted CRT device and compared them to patients with non LBBB and demonstrated that improved synchrony might translate into a reduction of ventricular arrhythmic events in LBBB patients. Ludwig et al.

(29) reported that patients with heart failure and LBBB, acute RVA pacing induces greater mechanical dyssynchrony and further impairs LV function. Sahebjam et al. (30) investigated the relationship between left atrial function and left ventricular dyssynchrony in heart failure patients. They revealed that left ventricular dyssynchrony was independently correlated with the deformity indices of the LA lateral wall.

The assessment of LV dyssynchrony during AMI has been performed using various echocardiographic techniques such as pulse wave TDI (31), color-coded TDI (2), speckle tracking (3), real time 3D echocardiography (32), and magnetic resonance imaging (MRI) (13). Although pulsed wave tissue Doppler is faster, the spatial resolution of this method is low and it has the disadvantage of comparing the measurements which are obtained from numerous different cycles. So heart rate variation is much more sensitive to the global motion and loading status of heart. Nevertheless it is reported that similar results with colored Doppler could be obtained (1). With the measurement of tissue displacement, in other words tissue tracking imaging, the visual and quantitative amount of tissue motion can be interpreted in millimeters-centimeters in a relatively easier way (33-36). As strain and strain rate imaging are not affected by impulsion-traction and translation, they may provide better differentiation of mechanical delay (37, 38). The frequently observed shortening following systole with these two methods causes the term "peaking time" to become more complicated. MRI is considered the gold standard for assessing cardiac shape and function, and other modalities such as real-time three-dimensional echocardiography (RT-3DE) (39, 40), closely correlate with MRI. Dyssynchrony using tissue Doppler images (TDI) during the acute phase of myocardial infarction is also considered a predictor of LV remodeling (3, 41) and although RT-3DE can be used to assess cardiac dyssynchrony (42), few reports have been published and the results do not coincide with those of TDI (43). The current study used TSI to detect dyssynchronous wall motion because of its ease of use and relatively good reproducibility. Starting point was set at the point of the opening of the aortic valves, and the end point was set at 200 milliseconds after the closure of the aortic valves, the beginning of the rapid filling period. In this way, the dyssynchronous motion within the systolic period was observed, whereas the latent influence of contractile motion during the isovolumic contraction period was excluded.

The patients in the current study had an initial EF of 37.7%. All had significant left ventricular systolic dysfunctions. In accordance with the outcomes of the studies in the literature, the patients of this study had distinct left ventricular dyssynchrony and their EF increased to 42.9% following percutaneous coronary intervention, while there was a significant decrease in left ventricular dyssynchrony. Different patterns were observed in pre PCI and post-PCI Ts values in anterior AMI and inferior AMI groups according to the infarction area. As expected in anterior AMI, the Ts values of the anterior segments in the post-PCI decreased. On the other hand, inferior segments were

596

affected by inferior infarction and the Ts values decreased in the post-PCI. The decrease of the Ts values in the basal septal segments is somewhat intriguing. This could imply that the septal wall of the LV is more vulnerable to develop dysynchrony in the occurrence of LV systolic dysfunction. Manka et al. (44) demonstrated that the regression of left ventricular dyssynchrony during healing of acute AMI. Zhang et al. (2) analyzed left ventricular systolic dyssynchrony in acute myocardial infarction patients with normal QRS durations. A total of 47 ST-segment elevation myocardial infarction patients were enrolled in the study and compared to the control group. Peak systolic velocity durations were found to be significantly longer in the AMI group. Again in this study, peak systolic velocity durations of anterior AMI patients was compared to those of inferior AMI patients and was found to be longer. A correlation between the degree of LV dyssynchrony and infarct area was detected. Nucifora et al. (45) investigated the effect of post-MI left ventricular functions on dyssynchronization in their study. Left ventricular dyssynchrony was found to be higher in patients with damaged left ventricle functions, large infarct areas, and in anterior AMI. However in these studies, there were no pre- or post-PCI comparisons made.

Finally, the accuracy and superiority of the TSI technique in the dyssynchrony of acute myocardial ischemia was demonstrated in a fewer number of clinical studies (46-49). The current study is important because myocardial dyssynchrony can be determined quantitatively in patients with AMI in the acute ischemic period and in the post-PCI period. The current results demonstrate the potential clinical advantages of using TSI in the detection of myocardial dyssynchrony changes in AMI patients in the early reperfusion period.

Study limitations

The present study has several limitations. Only longitudinal myocardial motion and dyssynchrony were examined in the present study, whereas radial and circumferential motions were not evaluated. The reproducibility and usefulness of TDI in the evaluation of mechanical dyssynchrony has been questioned recently (50). Recently the new methods such as speckle tracking imaging and three dimensional echocardiography were not used. The other limitation was the image quality and the artifacts. Its generation of a difference between the first hour admission and 12th hour admission in terms of myocardial damage, may effect the left ventricle dyssynchrony. The technique is highly dependent on the adequate training of operators and its reproducibility remains high in experienced laboratories. The measurements were done in the early period. The late results were not evaluated. In the near future, technique-related limitations can be overcome by the development of 2- and 3- dimensional or intra-cardiac strain echocardiography methods.

Conclusion

This study demonstrated that percutaneous coronary intervention greatly decreased LV dyssynchronization in the acute

period in patients with acute ST-segment elevation acute myocardial infarction. The deleterious effect of systolic asynchrony on global LV function may contribute to the acute remodeling process after AMI and one of the treatment targets in these patients should be LV dyssynchronization. There are many studies required in order to use this method routinely in LV dyssynchronization.

Conflict of interest: None declared.

Peer-review: Externally peer-reviewed.

Authorship contributions: Concept - S.İ., Ş.K.; Design - S.İ.; Supervision - Ş.K.; Resource - M.H.T., F.G., Ş.A.; Materials - E.M.B.; Data collection &/or processing - S.İ., H.D.; Analysis &/or interpretation - H.Ş.; Literature search - S.İ., Ş.K., M.H.T., Ş.A., F.G., E.M.B., H.D., H.Ş.; Writing - S.İ., M.H.T.; Critical review - S.İ., Ş.K., M.H.T., Ş.A., F.G., E.M.B., H.D., H.Ş.

References

- Sade LE, Özin B, Müderrisoğlu H. Echocardiographic evaluation of regional mechanical dyssynchrony and candidates for cardiac resynchronization therapy. Arch Turk Soc Cardiol 2005; 33: 413-22.
- Zhang Y, Chan AK, Yu CM, Lam WW, Yip GW, Fung WH, et al. Left ventricular systolic asynchrony after acute myocardial infarction in patients with narrow QRS complexes. Am Heart J 2005; 149: 497-503. [Crossref]
- Mollema SA, Liem SS, Suffoletto MS, Bleeker GB, van der Hoeven BL, van de Veire NR, et al. Left ventricular dyssynchrony acutely after myocardial infarction predicts left ventricular remodeling. J Am Coll Cardiol 2007; 50: 1532-40. [Crossref]
- Delgado V, Sitges M, Vidal B, Silva E, Azqueta M, Tolosana JM, et al. Assessment of left ventricular dyssynchrony by real-time three-dimensional echocardiography. Rev Esp Cardiol 2008: 61: 825-34. [Crossref]
- Gorcsan J 3rd, Kanzaki H, Bazaz R, Dohi K, Schwartzman D. Usefulness of echocardiographic tissue synchronization imaging to predict acute response to cardiac resynchronization therapy. Am J Cardiol 2004; 93: 1178-81. [Crossref]
- Gürel E, Tigen K. Selection of candidates for cardiac resynchronization therapy: late gadolinium enhanced cardiac magnetic resonance as a new and promising predictor of intraventricular dyssynchrony. Anadolu Kardiyol Derg 2011; 11: 263-8.
- Baldasseroni S, Opasich C, Gorini M, Lucci D, Marchionni N, Marini M, et al. Left bundle-branch block is associated with increased 1-year sudden and total mortality rate in 5517 outpatients with congestive heart failure: a report from the Italian network on congestive heart failure. Am Heart J 2002; 143: 398-405. [Crossref]
- 8. Fauchier L, Marie O, Casset-Senon D, Babuty D, Cosnay P, Fauchier JP. Interventricular and intraventricular dyssynchrony in idiopathic dilated cardiomyopathy: a prognostic study with fourier phase analysis of radionuclide angioscintigraphy. J Am Coll Cardiol 2002; 40: 2022-30. [Crossref]
- Yu CM, Lin H, Zhang Q, Sanderson JE. High prevalence of left ventricular systolic and diastolic asynchrony in patients with congestive heart failure and normal QRS duration. Heart 2003; 89: 54-60. [Crossref]
- 10. Bader H, Garrigue S, Lafitte S, Reuter S, Jais P, Haissaguerre M, et al. Intra-left ventricular electromechanical asynchrony. A new

- independent predictor of severe cardiac events in heart failure patients. J Am Coll Cardiol 2004; 43: 248-56. [Crossref]
- Penicka M, Bartunek J, Lang O, Medilek K, Tousek P, Vanderheyden M, et al. Severe left ventricular dyssynchrony is associated with poor prognosis in patients with moderate systolic heart failure undergoing coronary artery bypass grafting. J Am Coll Cardiol 2007; 50: 1315-23. [Crossref]
- Tian JW, Du GQ, Ren M, Sun LT, Leng XP, Su YX. Tissue synchronization imaging of myocardial dyssynchronicity of the left ventricle in patints with coronary artery disease. J Ultrasound Med 2007; 26: 893-7.
- Chang SA, Chang HJ, Choi SI, Chun EJ, Yoon YE, Kim HK, et al. Usefulness of left ventricular dyssynchrony after acute myocardial infarction, assessed by a tagging magnetic resonance image derived metric, as a determinant of ventricular remodeling. Am J Cardiol 2009; 104: 19-23. [Crossref]
- Nixon JV, Brown CN, Smitherman TC. Identification of transient and persistent segmental wall motion abnormalities in patients with unstable angina by two-dimensional echocardiography. Circulation 1982; 65: 1497-503. [Crossref]
- Gibson DG, Sanderson JE, Traill TA, Brown DJ, Goodwin JF. Regional left ventricular wall movement in hypertrophic cardiomyopathy. Br Heart J 1978; 40: 1327-33. [Crossref]
- Henein MY, Patel DJ, Fox KM, Gibson DG. Asynchronous left ventricular wall motion in unstable angina. Int J Cardiol 1997; 59: 37-45. [Crossref]
- 17. Antmann EM, Anbe DT, Armstrong PW, Bates ER, Green LA, Hand M, et al. ACC/AHA Guidelines for the management of patients with ST-elevation myocardial infarction: a report of the American College of Cardiology /American Heart Association task force on practice guidelines (Committee to revise the 1999 Guidelines for the management of patients with acute myocardial infarction) Circulation 2004; 110: 82-292.
- Yu CM, Zhang Q, Fung JW, Chan HC, Chan YS, Yip GW, et al. A novel tool to assess systolic asynchrony and identify responders of cardiac resynchronization therapy by tissue synchronization imaging. J Am Coll Cardiol 2005; 45: 677-84. [Crossref]
- Arita T, Sorescu GP, Schuler BT, Schmarkey LS, Merlino JD, Vinten-Johansen J, et al. Speckle-tracking strain echocardiography for detecting cardiac dyssynchrony in a canine model of dyssynchrony and heart failure. Am J Physiol Heart Circ Physiol 2007; 293: 735-42. [Crossref]
- Karakaş MF, Büyükkaya E, Kurt M, Karakaş E, Büyükkaya S, Akçay AB, et al. Assessment of left ventricular dyssynchrony in dipper and non-dipper hypertension. Blood Press 2013; 22: 144-50. [Crossref]
- Karakaş MF, Büyükkaya E, Kurt M, Çelik M, Karakaş E, Büyükkaya S, et al. Left ventricular dyssynchrony is an early manifestation of heart involvement in sickle cell anemia. Echocardiography 2013; 30: 521-6. [Crossref]
- Schiller NB, Shah PM, Crawford M, DeMaria A, Devereux R, Feigenbaum H, et al. Recommendations for quantitation of the left ventricule by two-dimensional echocardiography. American Soceity of Echocardiography Committee on Standards, Subcommittee on Quantitation of two-dimensional Echocardiograms. J Am Soc Ecocardiography 1989; 2: 358-67. [Crossref]
- Penicka M, Bartunek J, De Bruyne B, Vanderheyden M, Goethals M, De Zutter M, et al. Improvement of left ventricular function after cardiac resynchronization therapy is predicted by tissue Doppler imaging echocardiography. Circulation 2004; 109: 978-83. [Crossref]
- 24. Shin SH, Hung CL, Uno H, Hassanein AH, Verma A, Bourgoun M, et al. Mechanical dyssynchrony after myocardial infarction in

- patients with left ventricular dysfunction, heart failure, or both. Circulation 2010; 121: 1096-103. [Crossref]
- Cho GY, Song JK, Park WJ, Han SW, Choi SH, Doo YC, et al. Mechanical dyssynchrony assessed by tissue Doppler imaging is a powerful predictor of mortality in congestive heart failure with normal QRS duration. J Am Coll Cardiol 2005; 46: 2237-43. [Crossref]
- Cho GY, Kim HK, Kim YJ, Choi DJ, Sohn DW, Oh BH, et al. Electrical and mechanical dyssynchrony for prediction of cardiac events in patients with systolic heart failure. Heart 2010; 96: 1029-32. [Crossref]
- 27. Fauchier L, Marie O, Casset-Senon D, Babuty D, Cosnay P, Fauchier JP. Ventricular dyssynchrony and risk markers of ventricular arrhythmias in nonischemic dilated cardiomyopathy: a study with phase analysis of angioscintigraphy. Pacing Clin Electrophysiol 2003: 26: 352-6. [Crossref]
- Kutyifa V, Pouleur AC, Knappe D, Al-Ahmed A, Gibinski M, Wang PJ, et al. Dyssyncrony and the risk of ventricular arrhythmias. JACC Cardiovasc Imaging 2013; 6: 432-44. [Crossref]
- Ludwig DR, Tanaka H, Friehling M, Gorcsan J 3rd, Schwartzman D. Further deterioration of LV ejection fraction and mechanical synchrony during RV apical pacing in patients with heart failure and LBBB. J Cardiovasc Transl Res 2013; 6: 425-9. [Crossref]
- Sahebjam M, Zoroufian A, Sadeghian H, Roomi ZS, Sardari A, Mirzamani SS, et al. Relationship between left atrial function and size and level of left ventricular dyssyncrony in heart failure patients. Echocardiography 2013; 30: 772-7. [Crossref]
- Fahmy Elnoamany M, Mahfouz Badran H, Helmy Abo Elazm T, Shawky Abdelaziz E. Asynchrony of left ventricular systolic performance after the first acute myocardial infarction in patients with narrow QRS complexes: Doppler tissue imaging study. J Am Soc Echocardiogr 2006; 19: 1449-57. [Crossref]
- Maruyama Y, Masaki N, Yoshimoto N. Dyssynchrony during acute phase determined by real-time three dimensional echocardiography predicts reverse cardiac remodeling and improved cardiac function after reperfusion therapy. J Cardiol 2009; 54: 432-40. [Crossref]
- Sogaard P, Egeblad H, Kim WY, Jensen HK, Pedersen AK, Kristensen BO, et al. Tissue Doppler imaging predicts improved systolic performance and reversed left ventricular remodeling during longterm cardiac resynchronization therapy. J Am Coll Cardiol 2002; 40: 723-30. [Crossref]
- 34. Yu CM, Chau E, Sanderson JE, Fan K, Tang MO, Fung WH, et al. Tissue Doppler echocardiographic evidence of reverse remodeling and improved synchronicity by simultaneously delaying regional contraction after biventricular pacing therapy in heart failure. Circulation 2002; 105: 438-45. [Crossref]
- 35. Pan C, Hoffmann R, Kuhl H, Severin E, Franke A, Hanrath P. Tissue tracking allows rapid and accurate visual evaluation of left ventricular function. Eur J Echocardiogr 2001; 2: 197-202. [Crossref]
- Sade LE, Severyn DA, Kanzaki H, Dohi K, Gorcsan J 3rd. Secondgeneration tissue Doppler with angle-corrected color-coded wall displacement for quantitative assessment of regional left ventricular function. Am J Cardiol 2003; 92: 554-60. [Crossref]
- Breithardt OA, Stellbrink C, Herbots L, Claus P, Sinha AM, Bijnens B, et al. Cardiac resynchronization therapy can reverse abnormal myocardial strain distribution in patients with heart failure and left bundle branch block. J Am Coll Cardiol 2003; 42: 486-94. [Crossref]
- Sade LE, Severyn DA, Kanzaki H, Dohi K, Edelman K, Gorcsan J III.
 Comparative study of angle-corrected tissue velocity, displacement, strain and strain rate imaging to characterize mechanical dyssynchrony in left bundle branch block. Eur J Echocardiogr 2003; 4: 67. [Crossref]

- Arai K, Hozumi T, Matsumura Y, Sugioka K, Takemoto Y, Yamagishi H, et al. Accuracy of measurement of left ventricular volume and ejection fraction by new real-time three-dimensional echocardiography in patients with wall motion anomalies secondary to myocardial infarction. Am J Cardiol 2004; 94: 552-8. [Crossref]
- Nikitin NP, Constantin C, Loh PH, Ghosh J, Lukaschuk EI, Bennett A, et al. New generation 3-dimensional echocardiography for left ventricular volumetric and functional measurements: comparison with cardiac magnetic resonance. Eur J Echocardiography 2006; 7: 365-72. [Crossref]
- Zhang Y, Yip GW, Chan AK, Wang M, Lam WW, Fung JW, et al. Left ventricular systolic dyssynchrony is a predictor of cardiac remodeling after myocardial infarction. Am Heart J 2008; 156: 1124-32. [Crossref]
- Kapetanakis S, Kearney MT, Siva A, Gall N, Cooklin M, Monaghan MJ. Real-time three-dimensional echocardiography: a novel technique to quantify global left ventricular mechanical dyssynchrony. Circulation 2005; 112: 992-1000. [Crossref]
- Burgess MI, Jenkins C, Chan J, Marwick TH. Measurement of left ventricular dyssynchrony in patients with ischemic cardiomyopathy: a comparison of real-time three-dimensional and tissue Doppler echocardiography. Heart 2007; 93: 1191-6. [Crossref]
- 44. Manka R, Kozerka S, Rutz AK, Stoeck CT, Boesiger P, Schwitter J. A CMR study of the effects of tissue edema and necrosis on left ventricular dyssynchrony in acute myocardial infarction: implica-

- tions for cardiac resynchronization therapy. J Cardiovasc Magn Reson 2012; 14: 47. [Crossref]
- 45. Nucifora G, Bertini M, Marsan NA, Delgado V, Scholte JA, Ng AC, et al. Impact of left ventricular dyssynchrony early on left ventricular function after first acute myocardial infarction. Am J Cardiol 2010; 105: 306-11. [Crossref]
- Wang J, Abraham TP, Korinek J, Urheim S, McMahon EM, Belohlavek M. Delayed onset of subendocardial diastolic thinning at rest identifies hypoperfused myocardium. Circulation 2005; 111: 2943-50. [Crossref]
- Lin FC, Chang SH, Hsieh IC, Hung KC, Yeh SJ, See LC, et al. Time to peak velocity measurements by pulsed wave Doppler tissue imaging to quantify ischemia-related regional myocardial asynchrony. J Am Soc Echocardiogr 2004; 17: 299-306. [Crossref]
- Lee PW, Zhang Q, Yip GW, Wu L, Lam YY, Wu EB, et al. Left ventricular systolic and diastolic dyssynchrony in coronary artery disease with preserved ejection fraction. Clin Sci (Lond) 2009; 116: 521-9.
 [Crossref]
- Ng AC, Tran da T, Allman C, Vidaic J, Leung DY. Prognostic implications of left ventricular dyssynchrony early after non-ST elevation myocardial infarction without congestive heart failure. Eur Heart J 2010; 31: 298-308. [Crossref]
- 50 . Chung ES, Leon AR, Tavazzi L, Sun JP, Nihoyannopoulos P, Merlino J, et al. Results of the predictors of response to CRT (PROSPECT) trial. Circulation 2008; 117: 2608-16. [Crossref]