

The Effect of Precordial ST-Segment Depression in Inferior ST-Segment Elevation Myocardial Infarction on Left Ventricular Systolic Function :Two-Dimensional Echocardiographic Study.

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ABSTRACT

Background and Objective: The importance of precordial ST-segment depression in patients with early inferior ST-segment elevation myocardial infarction remain unclear. Many studies have reported that patients with precordial ST-segment depression appear to have large infarctions. The objectives of this study was to evaluate the effect of precordial ST-segment depression in patients with early inferior ST-segment elevation myocardial infarction on the left ventricular systolic function and left ventricular regional wall abnormalities.

Methods: Fifty eight patients with first inferior ST-segments elevation myocardial infarction (37 male, 21 female), their ages ranged from 30-91year,mean age 60.59 ± 11.21 who underwent thrombolysis in the Coronary Care Unit of Erbil Teaching Hospital for the period from August 2008 to August 2009 were included in this study. Two-dimensional echocardiography was performed in the first week of acute inferior myocardial infarction. Patients were classified according to the absence (group I,30 ,51.72%) or presence (group II ,28 , 48.28%) of precordial ST- segment depression .

Results: Group-II patients had a higher significant incidence of left ventricular systolic dysfunction (8,28.57%) than group I (2,6.67%), $P=0.027$,despite thrombolytic therapy (alteplase). Group-II patients had higher significant incidence of left ventricular regional wall abnormalities (12,42.9%) than group-I (3,10%), $P 0.00$,despite thrombolytic therapy (alteplase).

Conclusions: Early two-dimensional echocardiography is recommended for patients with inferior ST-segment elevation myocardial infarction associated with precordial ST-segment depression for the earliest detection of regional wall abnormalities and left ventricular systolic dysfunction.

Key words: Left ventricular systolic function .

INTRODUCTION:

Researchers have proposed that precordial ST segment depression is purely an electrocardiography (ECG) consequences of ST segment elevation in the inferior limb leads, without physiologic importance¹. Shah et al first reported that precordial ST segment depression was a marker for increased risk in patients with an inferior myocardial infarction .since then, its pathogenesis and prognostic importance have been heavily debated². Urgent two-dimensional echocardiography can play a crucial role in establishing the

diagnosis of acute myocardial infarction³. Studies based on echocardiographic diagnosis of heart failure after acute myocardial infarction ,suggest a prevalence rate of 2–3% of the population aged 45 years and older.⁴ Population-based studies suggest that the one-year mortality rate is of the order of 30–40%.⁵ The EuroHeart Failure study found that 9% of patients in the UK died during an admission, while almost a third of survivors were re-admitted within 12 weeks of discharge⁶. Eric D et al found that patients with acute inferior myocardial infarction with reciprocal ST-segment

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frequently developed congestive heart failure and cardiogenic shock than those with little or no precordial ST segment depression⁷. No previous studies done to study the effect of reciprocal ST-segment depression in the early phase of ST segment elevation myocardial infarction (STEMI) on the left ventricular systolic function and left ventricular regional wall abnormalities as compared to those without reciprocal ST segment depression. The objectives of this study were to evaluate the effect of precordial ST-segment depression on left ventricular systolic function and left ventricular regional wall abnormalities in the first week of inferior ST-segment elevation myocardial infarction.

PATIENTS AND METHODS:

We had studied 58 patients, 37 (53.79%) male and 21 (36.21%) female, their ages (30-91yr), mean age 60.59 ± 11.21 who had been diagnosed to have first inferior STEMI for the period from October 2008 to October 2009 in the Coronary Care Unit of Erbil Teaching Hospital. The inclusion criteria was those patients who had been admitted within 6 hour of onset of symptoms with chest pain lasting >30minutes accompanied by ECG signs ≥ 0.1 mv ST segment elevation in at least two leads (II,III, or aVF) associated with typical rise of troponin⁸. Precordial reciprocal ST-segment depression, defined as more than 1 mm, measured 80 ms after the J point in at least 2 of the leads V1-V4. Patients were classified according to the absence (group I) 30 patients (51.72%) or presence (group II) 28 patients (48.28%) of precordial ST-segment depression (Table-1). Patients who had left or right bundle branch block, previous angina pectoris, previous non STEMI, cardiogenic shock, severe persistent tachycardia, persistent serious arrhythmias, evidence of old infarction (significant Q waves in leads that had no ST segment elevation), ventricular paced rhythm, ventricular rhythm, ECG evidence

intraventricular conduction defects were excluded from this study. The presence or absence of the following risk factors were recorded: history of hypertension, diabetes mellitus, smoking, alcohol, family history of ischemic heart disease. Body mass index had been taken for all of the patient which is equal to weight of the patient/meter square of the length of the patient, obesity defined as $BMI \geq 30$ ⁹. Data on the hospital course were available for all patients. All patients received alteplase (tissue Plasminogen Activator) within 6 hours from the onset of symptoms of inferior STEMI, the dose given over 90 minutes bolus dose (15 mg) followed by 0.7 mg/kg body weight, but not exceeding 50 mg over 30 minutes and then 0.5 mg/kg body weight not exceeding 35 mg over 60 minutes¹⁰. Complete two-dimensional and Doppler echocardiography were performed within the first week of admission to the Coronary Care Unit in order to determine the left ventricular systolic function. All echocardiographic examinations were performed with an Philips Machine (Envisor USA), using a 2.5MHz transducer. Left ventricular ejection fraction was determined from apical and four chamber view using the Simpson's biplane formula: left ventricular systolic dysfunction (LVSD) defined as left ventricular ejection fraction $\leq 50\%$ ¹¹. Left ventricular segmental wall motion abnormalities was graded as follows: Normal at rest (1); hypokinetic-marked reduction in endocardial motion and systolic thickening (2), akinetic-virtual absence of inward motion and systolic thickening (3). and dyskinetic-paradoxical wall motion away from centre of the left ventricle in systole (4)¹². All possible transthoracic echocardiographic views were obtained for each patients including parasternal long axis, parasternal short axis, apical four chamber, two chamber and apical long axis view to check for the presence of hypokinesia, akinesia and dyskinesia.

Statistical methods: Chi square test used for the differences of proportion between

the different groups and p value < 0.05 considered as significant, SPSS version 16.0 (statistical package for social sciences) computer system by assistant of expert statistics.

RESULTS:

The total study group consisted of 58 patients with first inferior STEMI, 37 male (63.79%), 21 female (36.2 %) their ages ranged from 30-91 year, the mean ages was 60.59 SD \pm 11.21). The average age was nearly equal in group I & II, the proportion of men was higher in group I (21,70%) versus (16,57.1%) without statistical significant difference ,as shown in (Table1). There was no statistical significant difference in the incidence of obesity, hypertension, diabetes mellitus, current smoking, and family history of

Disease between the two groups, (Table1). Group-II patients had a higher significant incidence Of LVSD (8,28.57%) than group I (2,6.67%), $P=0.027$,despite thrombolytic therapy. as shown in (able-2), (Figure-1). Group-II patients had higher significant incidence of left ventricular regional abnormalities (12,42.9%) than group-I (3,10%), $P 0.00$, despite thrombolytic therapy, as shown in (Table-2), (Figure-2). Despite thrombolytic therapy basal inferior lateral akinesia was observed in 7 patients (25%) in group II versus 1 patients (3.3%) in group I ($P 0.02$), while there was no statistical significant difference in the incidence of basal inferior wall hypokinesia , apical dyskinesia between group II and group I as shown in (Tabe3), (Figure2).

Table 1: Baseline characteristics of two patient groups

Risk factors	Total patients with inferior STEMI 58		P value
	Group-I* No 30(51.72%)	Group-II** No28(48.28%)	
Men	21 (70%)	16 (57.1)	0.31
Age (yr)	60 \pm 11.2	61 \pm 11.2	0.54
Obesity	17 (56.67%)	18 (64.29%)	0.6
Hypertension	10 (33.33%)	11 (39.29%)	0.64
Diabetes mellitus	5(16.67%)	5 (17.86%)	0.9
Current smoking	9 (30%)	7 (25%)	0.7
Family history	7 (23.33%)	8 (28.57%)	0.64

*Group-1=Inferior ST-segment elevation myocardial infarction with out reciprocal ST-segment depression

**Group-2=Inferior ST-segment elevation myocardial infarction with reciprocal ST-segment depression

Table 2: Comparison of left ventricular systolic function between group-I & group-II patients.

Total patients with inferior STEMI No=58 (32.95%)			
	Group-I No=30(51.72%)	Group-II No=28(48.28%)	P value
LVSD in patients with inferior STEMI 10 (17.24%)	2 (6.67%)	8 (28.57%)	0.027*
LV regional WMA in patients with inferior STEMI 15(25.9%)	3 (10%)	12(42.9%)	0.00*

LVSD= Left ventricular systolic dysfunction, LV=left ventricle, WMA=wall motion abnormalities * p value significant

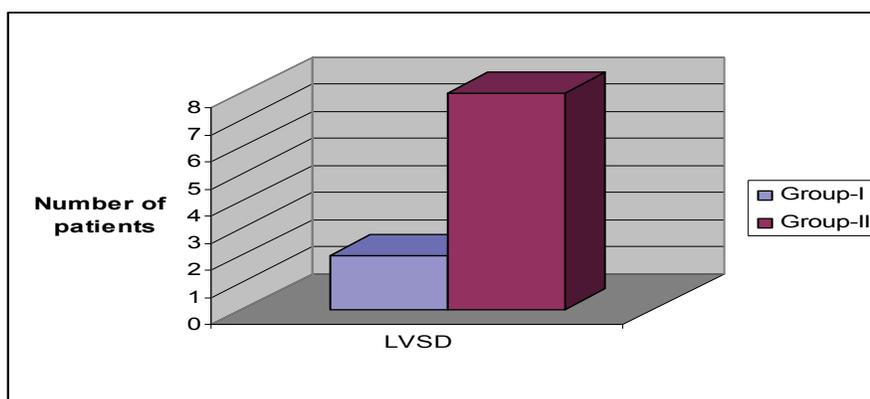


Figure 1: Comparison of the incidence of left ventricular systolic dysfunction(LVSD) between group I & II patients

Table 3: Comparison of grades of left ventricular regional wall abnormalities between group-I & II.

Left ventricular RWA	Site of left ventricular RWA	Group – I 30 (51.71%)		Group-II 28 (48.28%)		P
		No	%	No	%	
Hypokinesia	Basal inferior lateral	2	6.7	4	14.3	0.34
Dyskinesia	Apical	0	0	1	3.57	0.3
Akinesia	Basal inferior lateral	1	3.3	7	25	0.02*

RWA= Regional wall abnormalities, * = P value significant

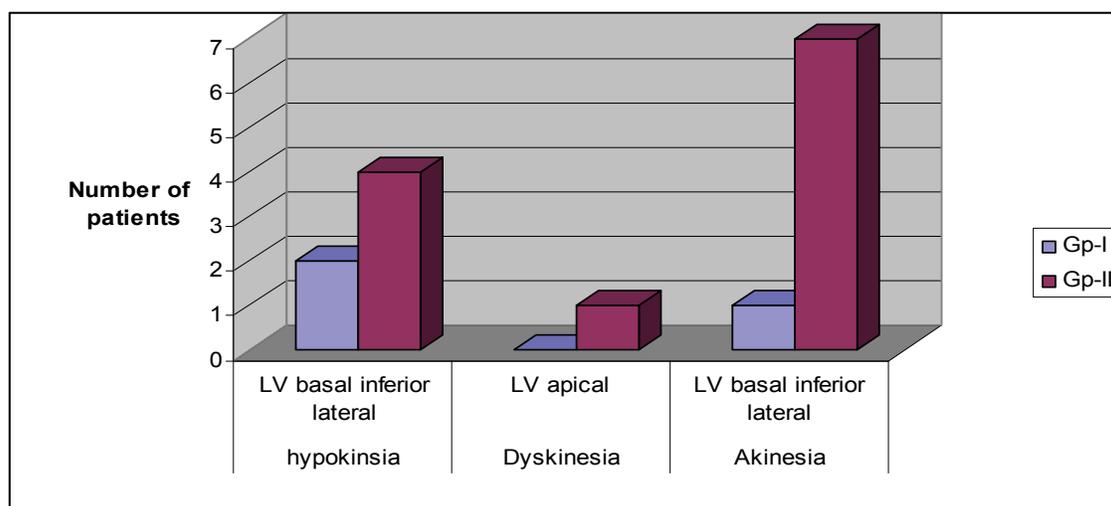


Figure 2: Comparison of left ventricular regional wall abnormalities between group-I & II patients

DISCUSSION:

Patients with inferior STEMI with precordial (reciprocal) ST-segment depression (group-II) had statistically significant incidence of LVSD 8 (28.57%) as compared with those without precordial (reciprocal) ST-segment depression (group-I) 2 (6.67%), $P=0.027$ as shown in (Table-2) & (Figure-1). Our results was similar to paterson et al study which found that left ventricular ejection fraction was significantly lower in patients who had ST segment depression in leads V1 to V3 or widespread ST segment depression than in those who had no ST segment⁷. Some studies have proposed that precordial ST segment depression during an inferior infarction signifies anterior wall ischemia and thus is a marker for left anterior descending coronary artery or multivessel disease^{13,14,15}. Tendra et al found that in patients with acute inferior myocardial infarction the presence and duration of ST-segment depression in leads v2 correlated with the presence of the left anterior descending coronary artery stenosis and depressed left ventricular ejection fraction¹⁶. We found a higher frequency rate of left ventricular regional wall abnormalities in patients with inferior STEMI who had reciprocal ST segment

(group-II) on 12-lead ECG 12 patients (42.9%) as compared with those patients with inferior STEMI without reciprocal ST-segment depression (group-I) 3 patients (10%), p value 0.00, as shown in (Table 2), Pierard et al found a higher statistical significant incidence of posterolateral wall akinesia in patient with acute inferior myocardial infarction with precordial reciprocal depression in the ECG versus those without reciprocal precordial ST depression, which is similar to our study¹. In conclusion early two-dimensional echocardiography is recommended for patients with inferior ST-segment elevation myocardial infarction associated with precordial ST-segment depression for the earliest detection of regional wall abnormalities and left ventricular systolic dysfunction. In order to provide appropriate early therapeutic decision before more deterioration in the left ventricular systolic function and a more conservative approach for those patients with inferior STEMI without precordial

STUDY LIMITATIONS:

ST-segment depression.

Our study had excluded those patients with severe persistent tachycardia, serious persistent arrhythmias because of difficulty

function in these situations. The results of this study should be confirmed in a larger number of unselected patients with inferior STEMI.

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