ORIGINAL ARTICLE

Predictors of No-Reflow in Patients undergoing Primary Percutaneous Coronary Intervention for Acute ST Elevation Myocardial Infarction at Tabba Heart Institute Karachi

AWAIS AHMAD NIZAMI, SHEHZAD AHMAD, BASHIR HANIF

¹Consultant Interventional Cardiologist, Cardiac Center. Bahawalpur.

²Assistant Professor of Cardiology and Director Cath Lab. Rai Medical College / RMC Teaching Hospital / Rai Medical Complex, Sargodha

³Medical Director and Chief of Cardiology, Tabba Heart Institute, Karachi. Pakistan Correspondence to Dr Awais Ahmad Nizami, E.mail:awaisnizami@hotmail.com, Cell: 03216831283

correspondence to Dr Awais Anmaŭ Nizami, E.mail.awaisnizami@notmail.com, Cell. 032 1663 126

ABSTRACT

Aim: To determine the predictors of no-reflow in patients undergoing Primary Percutaneous Coronary Intervention (PPCI) for acute ST elevation myocardial infarction (STEMI) at Tabba Heart Institute Karachi.

Study duration: The study was carried out from September 2020 to September 2021.

Methods: An observational study that was conducted at Tabba heart Institute Karachi. All patients with acute ST elevation myocardial infarction (STEMI) presenting within 12 hours of symptoms who undergoing primary PCI were included in the study. The patients were categorized Angiographically as no –reflow (with Thrombolysis in Myocardial Infarction (TIMI) <3 and reflow (TIMI-3) flow groups. The data regarding age and gender, type of MI, time from symptoms onset, co-morbid conditions like diabetes mellitus, hypertension, smoking, baseline Thrombolysis In Myocardial Infarction Trials (TIMI flow), infarct related artery (IRA), diameter of stent used, and the TIMI flow in IRA after the completion of procedure but before the use of intracoronary adenosine if needed, was recorded on a pre-designed proforma. The data was analyzed using SPSS 20.0.

Results: Total 296 patients who underwent primary PCI after acute MI were enrolled in the study. Mean age of the study population was 57±11 years. No reflow was present in 31.4% of patients. No reflow was more common in patients with age more than 60 years, females, symptom onset to primary PCI time of more than 6 hours, LAD infarct, and baseline TIMI flow < 2.

Conclusion: During primary PCI, no reflow occurs in a significant percentage of patients and is predictable by a number of factors which should be kept in mind to minimize the incidence of no reflow.

Key words: No reflow, acute myocardial infarction, primary PCI

INTRODUCTION

The standard care in an ST elevation Myocardial Infarction (STEMI) has been urgent reperfusion since long¹. No-reflow has been referred to as "sub optimal myocardial perfusion" of coronary artery segment with no angiographic evidence of obstruction². No-reflow is thought to be related to abnormal tissue perfusion, due to microvascular obstruction. The persistent no-reflow has shown and associated with poor clinical outcomes³. No- reflow leads, poor in-hospital and post discharge long term survival.⁴ It has been found to occur in 5 to 25 percent of cases⁵. In comparison to patients with adequate reflow after PCI, patients with no-reflow are more prone to develop complications like congestive heart failure early.⁶ The ongoing or persistent no-reflow is also linked to high mortality and an increased incidence of recurrent myocardial infarction.⁷ Patients with no-reflow also have relatively increased likelihood of early death (13% versus 6% p<0.003)⁸.

In the modern age of sophisticated interventional gadgets and techniques, no-reflow has still been shown to occur in 0.6% to 3.2% of PCI cases⁹. On Angiography the no-reflow has been referred to as "less than TIMI 3 flow" that occurs in 2% of all PCIs. TIMI grade-flow has been developed to ensure a standard reliable way of recording epicardial perfusion on coronary angiography. The ultimate finding of myocardial infarction reperfusion no-reflow would be consistent with a TIMI flow of less than 3. The findings from a study indicated that TIMI 2 flow is associated with a noreflow zone of substantial size; therefore, only TIMI 3 flow indicates reperfusion success¹⁰.

In view of the above background, it is of prime importance to identify the patients at risk as well as intra-procedural factors linked or predisposing to no-reflow after primary PCI to minimize the risk of complications as well as mortality in this population guiding management aiming to prevent, early intervention and active treatment strategy to prevent no-reflow occurring.

This study has been designed to determine the predictors of no-reflow in patients under-going primary PCI for acute myocardial infarction at Tabba Heart Institute Karachi.

Received on 12-12-2021 Accepted on 21-06-2022 The objective of the study was to determine the predictors of no-reflow on angiography, in patients undergoing primary PCI for acute ST Elevation myocardial infarction at Tabba Heart Institute Karachi.

MATERIALS AND METHODS

It was an observational study conducted at Tabba heart Institute Karachi after permission from Ethical Committee. All patients presenting with STEMI treated with PPCI were included in the study. They were divided into groups those who suffered no-reflow while others who did not have evidence of no-reflow. No-reflow was defined as: Angiographically TIMI <3

Grading system describes it as :

- 1. TIMI Grade 0 (no perfusion)
- 2. Grade 1:Contrast beyond obstruction but does not opacify the entire distal area
- 3. **Grade 2**: Contrast opacification distal to the obstruction however the rate of clearance is perceptibly slower
- 4. **Grade 3** (complete perfusion): Prompt antegrade flow distal to the obstruction1.1

Inclusion: All patients in all age groups with ST elevation myocardial infarction presenting within 12 hours of symptoms undergoing primary PCI were included in the study.

The Angiographic no –reflow (Thrombolysis in Myocardial Infarction (TIMI) <3 and reflow TIMI -3 flow was recorded.

Exclusion: Patients with presentation > 12 hours after symptoms, prior MI, prior CABG, cardiogenic shock at presentation, > 70% narrowing in the infarct related artery distal to the culprit lesion, or taking oral anti-coagulants for any reason were excluded from the study.

Convenience sampling was used to enroll the patients. All patients presenting in one year who fitted the inclusion criterion were included. The sample size of the study to detect 25.9% incidence of no reflow in primary PCI, with 95% power and error rate of 0.05 was 296 patients for a large STEMI population.

Data Collection: The data was collected on a pre-designed proforma. It included patient demographics including age and gender, type of MI, time from symptoms onset, co-morbid conditions like diabetes mellitus, hypertension, and smoking. Angiographic and procedural parameters that were included are:

baseline TIMI flow, infarct related artery, diameter of stent used, and the TIMI flow in IRA after the completion of procedure but before the use of intracoronary adenosine if needed. Coronary angiography images were read and evaluated by two cardiologists with more than 10 years' experience.

Data Analysis: Data was analyzed using SPSS 20. The categorical variables are presented as frequency and percentages. The continuous variables are presented as mean and standard deviation. The Chi square test was used to compare categorical variables and Student's *t*-test was used for comparisons between means (continuous variables). Significance will be assumed at p<0.05.

RESULTS

There were 296 patients who underwent primary PCI after acute MI were enrolled in the study from September 2020 to September 2021. Mean age of the study population was 57 ± 11 years. Study population comprised of 221(74.7%) males and 75(25.3%) females. No reflow was present in 93 patients (Fig-1). No reflow is more common in patients with age more than 60 years, females, symptom onset to primary PCI time of more than 6 hours, LAD infarct, baseline TIMI flow < 2, and baseline Killip Class > II. The patient characteristics in the two groups are shown in Table 1.



Fig 1: _Frequency of No reflow in patients undergoing primary PCI

Table 1: Characteristics of the patients undergoing primary PCI with and without no-reflow

	Total	Normal flow (203)		No reflow (93)		p Value
		No.	%	No.	%	
Age > 60 years	93 (31.4%)	03	1	90	96.8	<0.05
Female gender	77 (26%)	29	14.2	48	51.6	<0.05
Symptoms > 6hrs	157 (53%)	66	32.5	91	97.8	<0.05
DM	85 (28.7%)	61	30	24	25.8	>0.05
HTN	75 (25.3%)	55	27.1	20	21.5	>0.05
Smoking	175 (59.1%)	124	61.1	51	54.8	>0.05
Baseline TIMI flow < 2	76(25.7%)	28	13.8	48	51.6	
LAD infarction	176 (59.5%)	103	50.7	73	78.5	<0.05
LCX infarction	25 (8.4%)	21	10.3	04	4.3	
RCA infarction	84 (28.4%)	71	34.9	13	13.9	
Stent Diameter > 3 mm	91 (30.7%)	07	3.4	84	90.3	>0.05

DISCUSSION

In the recent era, primary PCI has emerged as the main stay of treatment of acute myocardial infarction. ¹²No reflow is a wellknown complication that occurs during primary PCI and is attributable to a number of factors including injury to the epithelium, neutrophil accumulation and distal thromboembolism.¹³No reflow has gained importance because it results in adverse short term and long term outcomes.

The main findings in our study are: (1) No reflow is encountered more frequently in people aged > 60 years. (2) female patients undergoing primary PCI are more likely to develop no reflow. (3) More time from symptom onset to primary PCI raises the chances of no reflow. (4) Baseline TIMI I/II flow raises the possibility of no reflow. (5) Primary PCI to LAD has more chances to encounter no reflow. (6) No reflow is more likely when larger diameter stent (>3 mm) is used in primary PCI.

Our findings are completely in accordance with the already existing data. Sabin et al^{5,13} in 2017 has revealed the incidence of no reflow in primary PCI to be 25.9%. Our study showed somewhat higher percentage of no reflow in primary PCI (31.4%). The finding of higher occurrence of no reflow in people aged > 60 years, females¹⁵ more time to reperfusion, poor baseline TIMI flow¹⁶ and LAD infarction are the same as in previous studies^{5,13,14} However we used a new parameter during our study which was stent diameter. It was found that stent diameter > 3 mm was associated with higher likelihood of no reflow. Similar reports have also been reported^{15,17}.

No reflow primarily affects endothelium and microvasculature, both negatively impact the presence of long-term systemic hypertension^{18,19} Hence, hypertension makes myocardial vasculature more susceptible and prone to No-reflow.

However this was not very significant amongst the two groups in our study. The reason behind differences could be ethnicity consistent with the published data, patients with no-reflow had worse outcomes²⁰.

The limitation of our study was that we used random sampling, which may have led to selection bias. Secondly the study is a single center study and the sample size was not very large.

CONCLUSION

The no-reflow is a known complication arising in primary PCI with various risk factors leading to morbidity and mortality. Prior Identification and management strategy significantly reduces the disease burden. The predictors in this study are almost similar to other ethnic groups in other parts of the world. They are easy to identify and we can use this factors for risk stratification. **Conflict of interest:** Nil

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