# Original ArticleFrequency of AcceleratedIdioventricular Rhythm in Patients withAcute ST Elevation Myocardial InfarctionReceiving Thrombolytic Therapy

Idioventricular Rhythm with Acute ST Elevation MI

Noshed Khan<sup>1</sup>, Samiullah Khan<sup>2</sup>, Naveed Danish<sup>4</sup>, Muhammad Niaz Khan<sup>5</sup>, Sadullah Shah<sup>1</sup> and Raza Muhammad<sup>3</sup>

## ABSTRACT

**Objective:** To determine the frequency of accelerated idioventricular rhythm in patients with acute ST elevation myocardial infarction receiving thrombolytic therapy.

Study Design: Descriptive study

**Place and Duration of Study:** This study was conducted at the Cardiology Department District Head Quarter Teaching Hospital (DHQ-TH) Bannu from February 2020 to August 2020.

**Materials and Methods:** A Total 372 patient of ST Segment elevated myocardial infraction (STEMI) aged 25 to 75 years of both gender were enrolled following written informed consent. Streptokinase was used as thrombolytic therapy and accelerated idioventricular rhythm (AIVR) was noted.

**Results:** In our study total 372 patients were enrolled with the mean age of  $53.8\pm11.9$  years. There were 53.2% males and 46.8% females. The most frequent risk factor was Hypertension in 83.9% patients, followed by Diabetes Mellitus 65.6%. Accelerated Idioventricular rhythm was present in 45.2% patients.

**Conclusion:** We observed that AIVR is a common reperfusion arrhythmia after thrombolysis with streptokinase **Key Words:** Myocardial infraction, reperfusion, ventricular arrhythemia

Citation of article: Frequency of Accelerated Idioventricular Rhythm in Patients with Acute ST Elevation Myocardial Infarction Receiving Thrombolytic Therapy. Med Forum 2021;32(11):33-37.

# **INTRODUCTION**

Myocardial infarction (MI) is the permanent loss of cardiac muscles caused by prolonged and an inadequate delivery of oxygen to the heart muscles.<sup>1</sup> It is caused by an imbalance in blood oxygen demand and supply owing to a variety of reasons, including plaque deposition and the formation of immovable thrombus in the coronary arteries that provide blood to the myocardium.<sup>2</sup>

In 2015, 15.9 million persons worldwide experienced a myocardial infarction, with more than 3 million having ST segment elevation and nearly 4 million having non-

<sup>5.</sup> Department of Cardiology, Hayatabad Medical Complex, Peshawar.

Correspondence: Dr. Samiullah Khan, Assistant Professor Cardiology, Bannu Medical College/ DHQ-TH, Bannu. Contact No: 03015151346 Email: drsamee@yahoo.com

Received:	June, 2021
Accepted:	August, 2021
Printed:	November, 2021

ST segment elevation MI.3,4 Severe chest discomfort extending to the lower jaw and shoulder is a frequent sign of MI. Preexisting coronary artery disease (CAD)<sub>6</sub>, high blood pressure, diabetes, smoking, poor nutrition, and lack of exercise<sup>7</sup> are all risk factors for MI. If the underlying reason is thrombus development in the coronary arteries, thrombolytic medicines are prescribed, which are administered intravenously to activate the fibrinolytic system in the blood, allowing plaques to be broken and blood flow to myocardial tissues to be restored.<sup>8,9</sup> Streptokinase, reteplase, alteplase, and tenecteplase are the most frequent thrombolytics utilized in acute ST segment raised MI (STEMI).<sup>10</sup> These medications are prescribed for individuals who have recently developed a STEMI and must be administered within 12 hours of the onset of symptoms in order to provide the most benefit to the patient.11

Resolution of ST segment elevation, reduction in chest discomfort, and the emergence of specific arrhythmias, particularly accelerated idioventricular rhythm, have all been proven to be valuable non-invasive markers for detecting coronary reperfusion in Acute MI.<sup>12-15</sup> Following fibrinolytic treatment, the AVIR is the most prevalent kind of arrhythmia, in the setting of STEMI.<sup>16</sup> Accelerated idioventricular rhythm, defined as a ventricular ectopic rhythm with more than 3 consecutive beats and a rate between 50 and 120 bpm,<sup>17-18</sup> is frequently observed during the reperfusion

33

<sup>&</sup>lt;sup>1.</sup> Department of Cardiology, DHO Bannu.

<sup>&</sup>lt;sup>2.</sup> Department of Cardiology / Medicine<sup>3</sup>, Bannu Medical College, DHQ-TH Bannu.

<sup>&</sup>lt;sup>4</sup> Department of Cardiology, DHQ Hospital Nowshera

#### Med. Forum, Vol. 32, No. 11

phase of acute myocardial infarction (AMI), and has therefore been proposed as a specific non-invasive marker for successful coronary artery reperfusion in the prethrombolytic and thrombolytic era.<sup>19-20</sup>

Acute intervention in the presence of reperfusion has not been demonstrated to enhance clinical outcomes, and coronary angiography should be postponed in instances when noninvasive reperfusion signs are of sufficient predictive value.<sup>21-22</sup> Emergency PTCA, on the other hand, is likely to be beneficial if coronary blood flow has not been restored. In Contrast if coronary blood flow has not been restored, emergency PTCA is likely to be helpful.<sup>23</sup>

## **MATERIALS AND METHODS**

Descriptive study conducted at the Cardiology department DHQ teaching hospital Bannu from 21<sup>st</sup> February 2020 to 20<sup>th</sup> August 2020.

**Sample Size:** Sample size is 372 patients, using 41% frequency of accelerated idioventricular rhythm in acute myocardial infarction receiving thrombolytic therapy, 95% confidence interval and 5% margin of error on WHO sample size calculator.<sup>17</sup>

Sampling Technique: Non probability consecutive sampling

### Sample Selection:

#### Inclusion criteria:

- Patients who presenting with chest pain within 12 hours' duration and ECG show ST segment elevation of more than 2 mm in chest leads or more than 1 mm elevation in limbs leads.
- Either gender.
- Age 25 years to 75 years.

#### **Exclusion criteria:**

- Those having contraindications to thrombolytic (Contraindications: CVA, active bleeding, suspected case of aortic dissection, malignant intracranial malignancy, head trauma)
- Previous myocardial infarction
  - Left bundle branch block.

Data Collection Procedure: This study was being carried out after the approval of the hospital Research Ethical Committee in the department of cardiology DHO-TH Bannu. The purpose and benefit of study was explained to patients and written inform consent was taken. Patients who fulfilled inclusion criteria were subjected to detail history and examination. Patients were monitored continually for 24 hours during and after infusion of thrombolytic (streptokinase) and appearance of accelerated idioventricular rhythm was noted. A twelve leads ECG was recorded by Fukuda Me C110 machine at standard paper speed of 25mm/second with 0.1 mm voltage representation standardization showing idioventicular rhythm. All information like age, gender, duration of disease, diabetes, hypertension and accelerated idioventricular rhythm was recorded using structured proforma.

**Data Analysis:** Data was analyzed using statistical package for social sciences version 16. Frequencies and percentages are calculated for categorical variables like gender, diabetes, and hypertension. Mean and standard deviation is calculated for numerical variables like age, duration of disease, accelerated idioventricular rhythm. Accelerated idioventricular rhythm is stratified among age, gender, diabetes, hypertension, and duration of disease in order to see effect modifiers. Post stratification chi square test is applied keeping p value < 0.05 as significant. Results are presented in tables and charts.

# RESULTS

In our study total 372 patients were enrolled with mean age of  $53.8\pm11.9$  years (28-75). There were 53.2% (n, 198) males and 46.8% (n, 174) female patients Table 1. Hypertension and Diabetes was present in 83.9% (n, 312) and 65.6% (n, 244) patients, respectively.

Frequency of accelerated Idioventricular rhythm was present in 45.2% (n, 168) patients as shown in Table 1.

**Table No.1. Baseline characteristics and Frequencies** 

Sr.	Variable	Freq-	%age
No		uency	
1.	Male	198	53.2%
2.	Female	174	46.8%
3.	Hypertensive	312	83.9%
4.	Normotensive	60	16.1%
5.	Diabetes	244	65.6%
6.	Non Diabetic	128	34.4%
7.	Idioventricular Rhythm	168	45.2%
	Present		
8.	Idioventricular Rhythm	204	54 804
	Absent	204	34.8%

Table No.2: Data stratification for age groups

			AIR		Total
			Yes	No	
	25 50	Count	75	66	141
Age	Age 25-50	% within Age group	53.2%	46.8%	100.0%
group	51-75 years	Count	93 138		231
		% within Age group	40.3%	59.7%	100.0%
Count			168	204	372
Total% withinAge group		45.2%	54.8%	100.0%	
p-value: 0.015 significant					

Age (p-value 0.015) and gender (p-value< 0.001), duration of MI (p-value 0.029) was significantly affecting the presence of AIR in the study subjects Table 2, 3 and 4.

Hypertension (p-value 0.978) and DM (p-value 0.255) was insignificantly associated with accelerated Idioventricular rhythm.

#### Table No.3: Data stratification for gender

			AIR	Total	
			Yes	No	
		Count	48	150	198
der	Male	% within Gender	24.2%	75.8%	100.0%
Gen	Female	Count	120	54	174
		% within Gender	69.0%	31.0%	100.0%
Count			168	204	372
Total % within Gender		45.2%	54.8%	100.0%	
p-value <0.001 significant					

Table No.4: Data stratif	ication for dura	ation of MI
	ATD	<b>T</b> - 4 - 1

			AIR		Total
			Yes	No	
		Count	27	15	42
	2-4	% within			
	hours	Duration of	64.3%	35.7%	100.0%
		MI			
		Count	39	45	84
W	4-6	% within			
of	hours	Duration of	46.4%	53.6%	100.0%
uo		MI			
ati		Count	48	79	127
m	6-8	% within			
П	hours	Duration of	37.8%	62.2%	100.0%
		MI			
		Count	54	65	119
	10-12	% within			
	hours	Duration of	45.4%	54.6%	100.0%
		MI			
		Count	168	204	372
Total		% within			
TOTAL		Duration of	45.2%	54.8%	100.0%
		MI			
p-value: 0.029 significant					

Table	No.5:	Data	stratification	for	diabetes
rabic	110.0.	Data	suameanon	101	ulancius

			AIR		Total
			Yes	No	
		Count	105	139	244
DM	Yes	% within Diabetes	43.0%	57.0%	100.0%
DM No		Count	63	65	128
	No	% within Diabetes	49.2%	50.8%	100.0%
Count		168	204	372	
Total% withinDiabetes		45.2%	54.8%	100.0%	
p-value: 0.255 not significant					

Table No.6:	Data	stratification	for	hypertension
1 abic 110.0.	Data	suamuanon	IUL	

			AIR		Total
			Yes	No	
		Count	141	171	312
Present	% within HTN	45.2%	54.8%	100.0%	
пти	HIN	Count	27	33	60
Absent		% within HTN	45.0%	55.0%	100.0%
Total		Count	168	204	372

	110		,
% within	45 204	54 804	100.00/
Hypertension	43.2%	54.0%	100.0%
p-value: 0.978 not significant			

# DISCUSSION

We observed in the present study that majority of patients with Acute STEMI were elderly.

A total 372 patients were enrolled with mean age of 53.8±11.9 years. This is in consistent with previous research<sup>24</sup> that that onset of this disease is mostly common in the older age. The predominant gender was male (53.2% males and 46.8% females) suffering acute STEMI. The most frequent risk factor observed was HTN and DM (83.9% and 65.6%, respectively) in our study. Khan S has reported even more male (Male:Female, 1:1.9) with Acute Myocardial infarction and Hypertension and Diabetes was the most frequent risk factor in the older age acute STEMI group patients in local population.<sup>25</sup> Khan A et al also confirmed almost similar results with mean age of  $59 \pm 10.8$ , (68%) male) and hypertension (n, 52), Smoking (n, 48) and diabetes (44) were the most frequent risk factor in elderly patients group.17

Frequency of AVIR in post thrombolytic patients with acute STEMI was observed in 45.2% (n, 168) in the present study. Khan et al reported AVIR 51% patients develop AIVR after thrombolytic therapy. Gressin et al<sup>20</sup> studied arrhythmias of ventricular origin during thrombolytic therapy administered for acute myocardial infarction using twenty-four-hour Holter monitoring in Ventricular patients treated with streptokinase arrhythmias were present in all patients. Tolerance was good (1 cardioversion for ventricular fibrillation). The incidence of AVIR was 90% with patent artery and 82% with non-patent artery. This frequency is almost double than our result which can be attributed to use of sophisticated holter monitoring as compared to random ECG monitoring done in our study. Wehren et al<sup>26</sup> found that AVIR was documented in 51% patients after thrombolytic therapy. These results are close to our results but it was done with small sample size of 110 patients.

In recent a study by Tatli et al<sup>27</sup> the incidence of AIVR in successfully thrombolysed patients was 73%. These all studies validate results of my study so my study can be used as reference study for taking AIVR as marker of reperfusion after thrombolytic therapy in the local population.

AIVR occurring in the first 6h were found significantly higher in patients with arterial patency and these arrhythmias were defined as non-invasive indicators of early coronary reperfusion.<sup>28</sup>

# CONCLUSION

To conclude, our study demonstrated the results that AIVR is the frequently recorded arrhythmia of

reperfusion during and or post thrombolysis with Streptokinase.

Author's Contribution:	
Concept & Design of Study:	Noshed Khan
Drafting:	Samiullah Khan, Naveed
	Danish
Data Analysis:	Muhammad Niaz Khan,
	Sadullah Shah, Raza
	Muhammad
Revisiting Critically:	Noshed Khan, Samiullah
	Khan
Final Approval of version:	Noshed Khan

**Conflict of Interest:** The study has no conflict of interest to declare by any author.

## REFERENCES

- 1. Thygesen K, Alpert JS, Jaffe AS, Chaitman BR, Bax JJ, Morrow DA, etal. Fourth universal definition of myocardial infarction 2018. JACC 2018;72(18):2231-64.
- 2. Salinas GL, Pascual M, Izco CF, Badano LP, Zamorano JL. Ischaemic heart disease: acute coronary syndrome. Echocardiol 2017:216.
- Vos T, Allen C, Arora M, Barber RM, Bhutta ZA, Brown Aetal. Global, regional, and national incidence, prevalence, and years lived with disability for 310 diseases and injuries, 1990–2015: a systematic analysis for the Global Burden of Disease Study 2015. The Lancet 2016;388 (10053):1545-602.
- 4. White HD, Chew DP. Acute myocardial infarction. The Lancet 2008;372;(9638):570-84.
- 5. Sabatine MS, Cannon CP. Approach to the patient with chest pain. Braunwald's Heart Disease: a textbook of cardiovascular medicine. 10th ed. Philadelphia, PA: Elsevier Saunders;2015.
- 6. Stone GW, Gao R, Kimura T, Kereiakes DJ, Ellis SG, Onuma Y et al. 1-year outcomes with the Absorb bioresorbable scaffold in patients with coronary artery disease: a patient-level, pooled meta-analysis. The Lancet 2016;387(10025): 1277-89.
- Schenck-Gustafsson K. Risk factors for cardiovascular disease in women. Maturitas 2009; 63(3):186-90.
- Malik AO, Abela O, Allenback G, Devabhaktuni S, Lui C, Singh A, et al. ST-segment elevation myocardial infarction, systems of care. An urgent need for policies to co-ordinate care in order to decrease in-hospital mortality. Int J Cardiol 2017; 240:82-6.
- 9. Collen D. Thrombolytic therapy. JTH 1997; 78(01):742-6.
- 10. Tough J, Berry L. Thrombolytic therapy in acute myocardial infarction. Nursing Standard (through 2013) 2005;19(37):55.

- 11. O'Gara PT, Kushner FG, Ascheim DD, Casey DE, Chung MK, De Lemos JA, et al. ACCF/AHA guideline for the management of ST-elevation myocardial infarction: a report of the American College of Cardiology Foundation/American Heart Association Task Force on Practice Guidelines. JACC 2013;61(4):e78-140.
- 12. Marriott JCJL, Menendez MM. A-V dissociation revisited. ProgCardiovasc Dis 1966;8: 522–538.
- 13. Krucoff MW, Croll MA, Pope JE, et al. Continuously updated 12-lead ST segment recovery analysis for myocardial infarct artery patency assessment and its correlation with multiple simultaneous early angiographic observations. Am J Cardiol 1993;71:145–51.
- 14. Klootwijk P, Langer A, Meij S, et al. Non-invasive prediction of reperfusion and coronary artery patency by continuous ST segment monitoring in the GUSTO-1 trial. Eur Heart J 1996;17:689–98.
- CaliV RM, O'Neill W, Stack RS, et al. Failure of simple clinical measurements to predict reperfusion status after intravenous thrombolysis. Ann Int Med 1988;108: 658–62.
- 16. Doevendans PA, Gorgels AP, Van der Zee R, et al. Electrocardiographic diagnosis of reperfusion during thrombolytic therapy in acute myocardial infarction. Am J Cardiol 1995;75:1206–10.
- 17. Khan A, Nadeem S, Kokane H, Thummar A, Lokhandwala Y, Mahajan AU, et al. Is accelerated idioventricular rhythm a good marker for reperfusion after streptokinase? Ind Heart J 2016; 68(3):302-5.
- Grimm W, Hoffmann J, Maisch B. Akzelerierteridioventrikulärer rhythmus. Z Kardiol 1994; 83:898–907.
- 19. Goldberg S, Greenspon AJ, Urban PL, *et al* Reperfusion arrhythmias: A marker of restoration of antegrade flow during intracoronary thrombolysis for acute myocardial infarction. Am Heart J 1983;105: 26–31.
- 20. Gressin V, Louvard Y, Pezzano M, *et al.* Holter recording of ventricular arrhythmias during intravenous thrombolysis for acute myocardial infarction. Am J Cardiol 1992;69:152–159.
- 21. Simoons ML, Arnold AER, Betriu A, et al. European Cooperative Study Group for recombinant tissue-type plasminogen activator (rTPA). Thrombolysis with tissue plasminogen activator in acute myocardial infarction: no additional benefit from immediate percutaneous coronary angioplasty. Lancet 1988;i:197–203.
- 22. TIMI research group. Immediate vs delayed catheterization and angioplasty following thrombolytic therapy for acute myocardial infarction. TIMI 2A results. JAMA 1988;260: 2849–58.

- 23. Grines CL, Browne KF, Marco J, et al. Primary angioplasty in myocardial infarction study group: a comparison of immediate angioplasty with thrombolytic therapy for acute myocardial infarction. N Engl J Med 1993;328:680–4.
- 24. Roth GA, Johnson C, Abajobir A, Abd-Allah F, Abera SF, AbyuG,etal. Global, regional, and national burden of cardiovascular diseases for 10 causes, 1990 to 2015. J Am Coll Cardiol 2017; 4:70(1):1-25.
- 25. Khan S, Asghar Khan M, Khan MN, Shah I, Hassan M ul. Comparison of Risk Factors Profile in Patients Below and Above Forty Years of Age Presenting With Acute Myocardial Infarction. J Postgrad Med Inst 2013;27(4).
- 26. Khan A, Nadeem S, Kokane H, Thummar A, Lokhandwala Y, Mahajan AU, et al. Is accelerated idioventricular rhythm a good marker for reperfusion after streptokinase? Ind Heart J 2016:1;68(3):302-5.
- 27. Tatli E, Alicik G, Buturak A, Yilmaztepe M, Aktoz M. Arrhythmias following revascularization procedures in the course of acute myocardial infarction: are they indicators of reperfusion or ongoing ischemia? Sci World J 2013.
- Cercek B, Horvat M. Arrhythmias with brief, highdose intravenous streptokinase infusion in acute myocardial infarction. Eur Heart J 1985; 109–113.