RESULTS

Results

Basic characteristics of the study group

The study group consists of **60** patients, with an age ranging from 39 to 70 years. It includes **53** males (88.3%) and **7** females (11.6%). **41** (68.3%) patients are hypertensive, **29** (48.3%) patients are diabetics and **45** (75%) patients are smokers, **33** (55%) patients are dyslipidemic and **14** (23.3%) patients have positive family history for coronary artery disease.

Table (3) : Basic characteristics of the whole study group:

Number of patients		60			
Age in years		Ranging (39 – 70 years)			
-mean age		55 ± 15 years			
Sex					
-Male	number (%)	53 (88.3 %)			
-Female	number (%)	7 (11.6%)			
Hypertensives	number (%)	41 (68.3%)			
Diabetics	number (%)	29 (48.3%)			
Smokers	number (%)	45 (75%)			
Dyslipidemics	number (%)	33 (55%)			
Positive FH	number (%)	14 (23.3%)			
Type of MI					
-Anterior		39 (56%)			
-Non-anterior		21 (35%)			
Time to reperfu	sion therapy	Ranging (1.30 – 10.30 hrs)			

According to the mode of reperfusion the study group was classified into:

GROUP I, it includes 30 patients who received streptokinase as a treating modality.

GROUP II, it includes 30 patients who underwent primary intervention as a treating modality.

Both groups were comparable as regard age ,sex ,risk factors for coronary artery disease, location of myocardial infarction and time to intervention .

Table (): Comparison of the clinical data in both groups

Clinical data	Group I		Group II		P-value
	Thrombolysis/30		Angiographic/30		
1-Age	Rang	ge (39-70)	Range(49-70)		>0.05
mean age	55 ±	15 years	60 ± 10 years		
2-Sex					
males	26	86.6%	27	90%	>0.05
females	4	13.3%	3	10%	
3-Risk factors					
Smoking	20	66.6%	25	83%	>0.05
Hypertension	21	70%	20	66.6%	>0.05
Diabetes	14	46.6%	15	50%	>0.05
Family history	6	20.0%	8	26.6%	>0.05
Dyslipidemia	14	46.6%	19	63.3%	>0.05
4-Previous IHD	11	36.6%	16	53.3%	>0.05
5-Type of MI					
-Anterior	2	66.6%	19	63.3%	>0.05

-Non-anterior	10 33.3%	11 36.6%	>0.05
6-therapy	Streptokinase	Primary PCI	
7-Time to therapy	Range	Range	
Pain to hospital	(1-7 hrs)	(1-10 hrs)	>0.05
Pain to therapy	(2-8 hrs)	(1.30-10.30hrs)	>0.05
Hospital delay	1.3 ± 0.9 hours	1.2 ± 0.7 hours	>0.05
8-Cardiac enzymes			
-Total CK	1997 ± 224	1229 ± 376	>0.05
-Ck-mb	122 ±187	101 ± 175	>0.05
-Peak time	14.5 ± 5	10.05 ± 3	>0.05

IN-HOSPITAL COMPLICATIONS

During hospitalization (after therapy) the incidence of left ventricular dysfunction diagnosed clinically was more in the thrombolytic group than in the angioplasty group, 7/30 (23.3%) VS 5/30 (16.6%)) respectively, P<0.05).

The incidence of life threatening arrythmias VF or VT that needed treatment were higher in the thrombolytic group than in the angioplasty group 9/30 (30%) VS 15/30 (16.6%) respectively, P<0.05.

On the other hand, the incidence of recurrent ischemia were significantly higher in the thrombolytic group than in the angioplasty group 11/30(36.6%)~&~9/30~(16.6%) respectively,P<0.05 and the incidence of infarctions were significantly higher in the thrombolytic group than in the angioplasty group 2/30~(6.6%)~&~1/30~(3.3%) respectively,P<0.05 .

The mortality rate during hospitalization in the angioplasty group was highly significantly less than that in the thrombolysis group 1/30 (3.3%) VS 4/30 (13.3%) respectively, P<0.05.

Table (): In-hospital complications in both groups

Clinical	Group 1		Gr	oup 2	P-value
complications	thrombolysis/30		Angioplasty/30		
1-Clinical LV	7	23.3%	5	16.6 %	< 0.05
dysfunction					
2-Recurrent	11	36.6%	9	16.6%	< 0.05
ischemia					
3-Infarction	2	6.6 %	1	3.3%	< 0.05
4-Life threating	9	30%	5	16.6%	< 0.05
arrythmias					
5-Bleedings	3	10%	2	6.6%	>0.05
-hematuria	2	6.6%			
-hematoma at	1	3.3%	2	6.6%	
puncture site					
6-Mortality	4	13.3%	1	3.3%	<0.05

PRE-DISCHARGE ANGIOGRAPHIC DATA

1-The extent of CAD was almost similar in both groups; more than one vessel affection was present in 19/30 (63.3%) of patient in both groups, single vessel disease was present in 11/30 (36.6%) of patients in both groups, P > 0.05.

- 2-Totally occluded IRA (100% stenosis) was found in a significantly higher incidence in the thrombolysis group than in the angioplasty group 12/30 (41%) VS 3/30 (10%) respectively, P<0.05.
- 3-The mean residual stenosis of the IRA was significantly higher in the thrombolysis group; $(91.56 \pm 16.7\% \text{ VS } 37 \pm 28.7\%)$ in the angioplasty group, P<0.05.
- 4-The achieved TIMI grade 3 flow (complete reperfusion) was significantlly higher in the angioplasty group than in the thrombolysis group 8/30 (83.3%) VS 20/30 (26.6%) respectively while the presence of TIMI flow 0-1 was significantly higher in the thrombolysis group 12/30 (40%) VS 3/30 (10%) respectively,P<0.05.
- 5-The patency rate was significantly higher in the angioplasty group than in the thrombolysis (90% VS 59%) respectively,P<0.05.

Table (): Angiographic data in both groups

Clinical data	Group 1		Group 2		P-value
	thrombolysis		angioplasty		
1-Extent of CAD					
-Single vessel	11	36.6%	11	36.6%	>0.05
-2 vessels	8	26.6%	12	40%	< 0.05
-3 vessels	11	36.6%	7	30%	< 0.05
->1 vessel	19	63.3%	19	63.3%	>0.05
2- IRA					
-LAD	20	66.6%	19	63.3%	>0.05

-Non-LAD	10	33.3%	11	36.6%	>0.05
3-Residual stenosis	mean(91.6%)		mea	n(37%)	< 0.05
of IRA:					
-Total occlusion	12	41%	3	10%	< 0.05
-< 50%	1	3.3%	25	83.3%	< 0.05
-> 50 %	29	96.6%	5	16.6%	< 0.05
4-TIMI flow in					
IRA					
-0 – 1	12	40%	3	10%	< 0.05
-2	10	30.3%	1	3.3%	< 0.05
-3	8	26.6%	26	83.3%	< 0.05
5-Patency rate	18	59%	27	90%	< 0.05

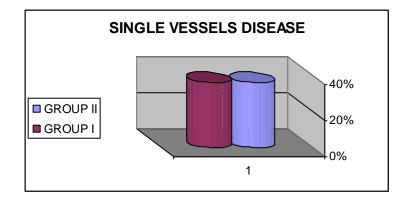
Left ventricular ejection fraction (LVEF) measured by Echocardiogram

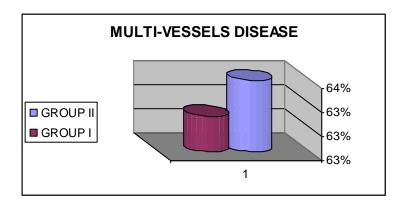
LVEF measured by echocardiogram at discharge was significantly better in the angioplasty group (50.2 + 9.7%) than that in the thrombolysis group (40.88 + 7.5%), P<0.05.

Table (7): (LVEF) measured by Echocardiogram

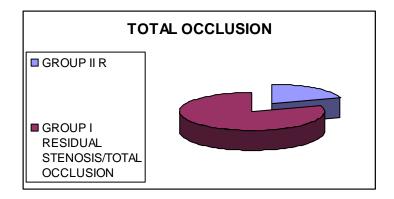
(LVEF)	Group 1	Group 2	P-value
	thrombolysis	angioplasty	
Echocardiogram	$40.88 \pm 7.5\%$	$50.2 \pm 9.7\%$)	< 0.05

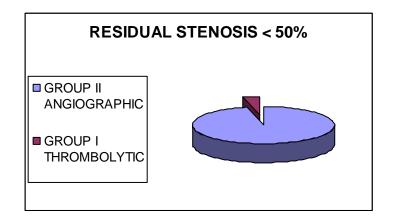
ANGIOGRAPHIC DATA OF THE VESSELS AFFECTED

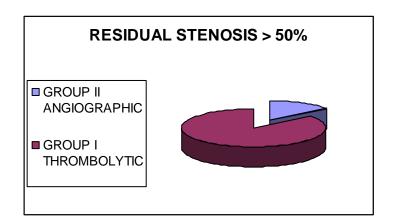




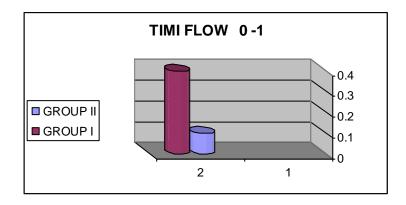
RESIDUAL STENOSIS

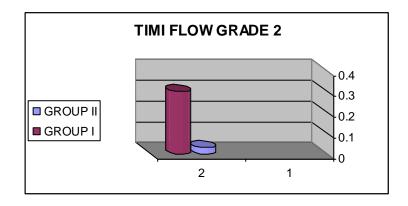


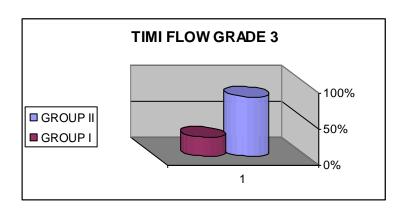




TIMI FLOW GRADE IN THE IRA



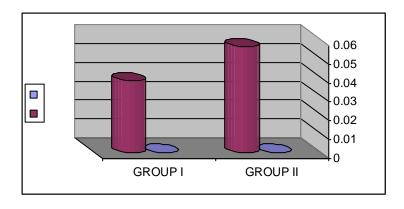




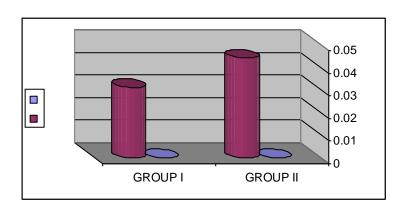
Time /hours

TIME TO TREATMENT

PAIN TO HOSPITAL

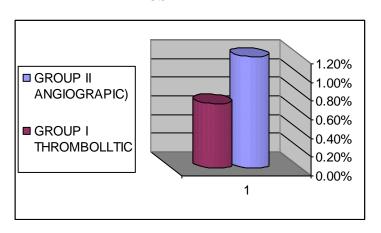


PAIN TO THERAPY



Time/hours

HOSPITAL DELAY



Time/hours

COMPLICATIONS

