Abstract

Is cardiac rehabilitation after PCI as effective as CABG? The first experience from the eastern mediterranean region cardiac rehabilitation registry

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OriginalArticle

BACKGROUND: The effectiveness of cardiac rehabilitation (CR) programs following either percutaneous coronary intervention (PCI) or coronary artery bypass grafting (CABG) has been separately studied. Few studies have compared the effects of similar CR programs between PCI and CABG. This study aimed to compare the effects of CR in patients recruited following either PCI or CABG on coronary heart disease risk factors, psychological variables, and functional capacity.

METHODS: For this retrospective study, the documents of the CR program registry of the Isfahan Cardiovascular Research Institute were reviewed from 2008 to 2021. Patients with ischemic heart disease undergoing PCI or CABG were enrolled in an 8-week exercise-based cardiac rehabilitation program. Demographics, smoking status, clinical data, echocardiographic parameters, laboratory data, functional capacity, and psychological status were assessed.

RESULTS: Patients who underwent CABG (n=557) were more likely to be referred to CR than those who underwent PCI (n=440). All variables changed significantly after the CR program compared to their baseline value in both the PCI and CABG groups. However, low-density lipoprotein and total cholesterol levels, peak systolic blood pressure, and resting and peak diastolic blood pressure did not change in any of the groups, and fasting blood sugar (p=0.01) and triglyceride (TG) (p=0.01) levels significantly decreased only in the PCI group. Between-group comparisons indicated that after adjustment, no significantly reduced in the PCI group (p=0.01).

CONCLUSION: The CR program was equally effective in patients who underwent either PCI or CABG.

Keywords: Cardiac Rehabilitation; Percutaneous Coronary Intervention; Coronary Artery Bypass Grafting; Coronary Heart Diseases; Psychological Factors; Risk Factor

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Introduction

Cardiovascular diseases, which are the leading cause of death globally, have several modifiable risk factors^{1,2} and can be managed and intervened upon by comprehensive exercise-based rehabilitation programs³. Cardiac rehabilitation (CR) programs have been introduced to patients following coronary events to facilitate lifestyle changes⁴⁻⁶. Indeed, CR has significant positive effects on the functional capacity, lipid profile, glycemic control, echocardiographic indexes, smoking behavior, and blood pressure of patients⁶⁻⁸. These programs can also enhance the quality of life, modify psychological factors, and reduce mortality and readmission rates^{9,10}. Although CR is highly recommended for all patients with coronary artery disease, referral to and participation in CR is globally low¹¹⁻¹³.

There are extensive studies on the effectiveness of CR after coronary artery bypass grafting (CABG), leading to an accumulation of evidence in favor of CR following this intervention⁴. Revascularization in patients with coronary artery diseases is also treated with less invasive procedures like percutaneous coronary intervention (PCI). Numerous studies have evaluated the impacts of CR in PCI only¹⁰⁻¹⁴; however, there is a lack of evidence from the Middle-Eastern region in this regard. Furthermore, no study has ever compared the effectiveness of CR after PCI to that of CABG. Therefore, the authors aimed to compare the impact of phase-II comprehensive CR after PCI vs. CABG on coronary heart disease risk factors, psychological variables, and functional capacity of the CR registry in the Eastern Mediterranean region. The authors hypothesize that if the value of CR after PCI is not more than that of CABG, it is not less than that, and both PCI and CABG patients will benefit from CR to an equal magnitude.

Methods

Study design

For this retrospective study, the CR program registry of the Cardiac Rehabilitation Research Center of the Cardiovascular Research Institute (a WHOcollaborating center in EMRO) was searched and reviewed from January 2008 to December 2021. All patients with ischemic heart disease who were admitted for either PCI or CABG were advised to participate in this hospital-based CR program. Before being discharged, an invitation card was given to them, which needed to be validated by their cardiologist or surgeon before participating in the program. The inclusion criteria were all registered patients who had undergone either PCI or CABG for the first time, completed the CR program as scheduled, and answered all the questionnaires. The exclusion criteria included the following: patients with serious medical conditions (e.g., cerebral vascular attacks, chronic kidney disease, cirrhosis, and chronic obstructive sleep apnea), patients who couldn't tolerate physical activity sessions, > 20% missing data in the medical documents or questionnaires, a previous history of PCI or CABG, and missing two or more CR program sessions.

Cardiac rehabilitation program:

CR was recommended to every patient with any indications of CR. This 8-week exercise-based CR program included both physical exercise and educational sessions. The physical exercise sessions were offered three times a week for eight weeks (24 sessions in total) and supervised by a trained sports physician. The eight lecture-based educational sessions for controlling stress, anxiety, and depression, as well as for quitting smoking, were led by a trained psychologist. The sessions on following a healthy lifestyle and nutrition plan were led by a trained dietician. The patients were contacted regularly before their sessions by the center secretary and reminded of the scheduled classes.

Assessments

A checklist of demographic variables (age and sex), smoking status (current, former, and never), physical activity level, laboratory data, cardiac function test results, and psychological status was used at the time of registration (within one week before starting the program), and was repeated within one week of completing the program.

To assess the physical activity level, the Persian validated long-form version of the International Physical Activity Questionnaires (IPAQ) was used¹⁵. IPAQ is a 7-day recall questionnaire that measures time spent per week on vigorous activity, moderate activity, and walking. Briefly, IPAQ assesses physical activity undertaken across a comprehensive set of domains (work, transportation, housework, and leisure time). Activity is then calculated as the total

time (in minutes) spent in three activity categories. The total time in each category is then weighted by a Metabolic Equivalent of Tasks (METs). According to the reported METs, subjects were categorized into three levels of activity: walking, moderate, and vigorous¹⁵.

Fasting blood samples were obtained before starting and after completing the program. All the samples were taken in the central laboratory at the center by the same team. Fasting blood glucose (FBS) and a lipid profile (triglyceride (TG), total cholesterol (TC), high-density lipoprotein cholesterol (HDL-C), and low-density lipoprotein cholesterol (LDL-C)) were recorded.

Echocardiography was scheduled for all patients before starting and after completing the CR. All echocardiographies were performed in the left lateral decubitus position with the Philips IE33 ultrasound machine and interpreted by an echocardiologist under standard protocols to obtain the left ventricular ejection fraction (EF)¹⁶.

The computer-controlled treadmill exercise test (Stress Test System, AST-3000, AVECINA Company, Iran) was used to evaluate functional capacity. The resting heart rate (HR), systolic, and diastolic blood pressure (SBP and DBP, respectively) were measured manually under the standard protocol before the exercise by an experienced exercise test room nurse. The intensity of the exercise test was scheduled with the graded multi-stage maximal symptomlimited Bruce protocol^{17,18}, which was continued until physical exhaustion or serious signs/symptoms occurred. The HR, SBP, and DBP were measured once at every stage, at peak exercise, and twice during the recovery phase. After completion, test duration, cardiorespiratory function in METs (derived from the walking speed and slope), and electrocardiography were extracted from the program. The final result of the exercise test was interpreted by a cardiologist and categorized as positive, negative, or undetermined/ unidentifiable.

To assess psychological status, the authors used the validated Persian versions of the questionnaires for anxiety¹⁹, depression²⁰, general quality of life (QoL)²¹, and health-related quality of life (HR-QoL)²².

Anxiety was evaluated with the 20-item Zung's self-rating anxiety scale (SAS) questionnaire²³ with scores ranging from normal to mild (20-44),

moderate (45-59), severe (60-74), and very severe (75 and above).

The level and score of depression were assessed using the Beck depression inventory (second edition) questionnaire (BDI-II), which consists of 21 questions²⁰ with scores of low (0-10), mild (11-16), moderate (17-30), and high (31 and above).

The SF-36 questionnaire was utilized by the authors to evaluate the general aspects of QoL²¹. This questionnaire has two general domains, namely physical and mental health, each with four subdomains. HR-QoL in cardiac disease was evaluated using the 27-item MacNew questionnaire²² with questions classified into physical, emotional, and social domains.

Statistical Analysis

The authors carried out all analyses with IBM SPSS software version 20.0. Categorical variables are expressed as numbers and percentages, while quantitative variables are expressed as mean and standard deviation. The Kolmogorov-Smirnov test was used to verify the normality assumption. For quantitative variables, a baseline measurement was assessed by an independent t-test or Mann-Whitney test (if the normality assumption was not met). Categorical variables were compared using the Chi-square test. Bonferroni correction was applied to determine the significance of any differences. Within-group comparisons were assessed by paired t-tests for normally distributed variables or Wilcoxon for non-normally distributed variables. tests Analysis of covariance (ANCOVA) was used to evaluate between-group comparisons. Variables that were significantly different at baseline or were confounders were also adjusted in ANCOVA. If the heterogeneity of variance was not met, a logarithmic transformation was applied. P-values < 0.05 (twotailed) are considered statistically significant.

Results

The CR program was conducted more frequently following CABG than PCI (n=557 vs. n=440 patients, respectively). Among these patients, male participation was higher than female, but there was no significant difference between the two genders (426 (76.48%) in CABG and 316 (71.81%) in PCI, p=0.1). The authors found that CABG patients were significantly older than the PCI patients (58.94 \pm 8.85

vs. 57.72 \pm 9.79 years, p= 0.02). 111 documents were excluded from the secondary analysis due to missing data after CR assessments.

CABG participants had significantly higher LDL-C (p<0.0001) and TC (p<0.0001) levels compared to PCI. However, the mean EF (p=0.01), exercise test METs (p=0.019), anxiety (p<0.0001), and depression (p<0.0001) scores were significantly higher in PCI (Table 1).

As shown in Table 2, all variables underwent significant changes after the CR program compared to their baseline value in either the PCI or CABG group. However, LDL-C and TC levels, peak SBP, and resting and peak DBP remained unchanged in all groups. Only in the PCI group did FBS (p=0.01) and TG (p=0.01) levels significantly decrease.

Between-group comparisons, after full adjustment, indicated no significant change after the CR program between the PCI and CABG groups, except for TG (Table 3). The authors observed a significant reduction in TG after the CR program in patients with PCI compared to those with CABG.

Discussion

It is generally believed that CR should be recommended to all patients with cardiovascular disease as a secondary prevention strategy²⁴. However, the outcomes of CR have not been compared between PCI and CABG patients in a comprehensive study from an advanced CR center in the Eastern Mediterranean region. The authors' results suggest that both PCI and CABG patients benefited similarly from CR, as the outcomes were not significantly different between the two groups in most examined variables. These data indicate that CR is a highly effective secondary prevention strategy in coronary artery disease patients, and its priority after PCI is as equal as after CABG.

Although the goal of CR is to educate patients about the harmful effects of smoking on the heart, its efficacy is not comparable to explicit smoking cessation programs in addiction treatment centers. More than half of the Portuguese CR participants quit smoking in the follow-up evaluations, and the authors have suggested that CR is a great opportunity to educate patients and emphasize the importance of smoking cessation. In this study, the distribution of smoking status changed significantly before and after CR in each group²⁵. This conclusion is in agreement with other studies, but without significant differences between PCI and CABG^{26,27}.

The positive effect of CR on functional capacity after PCI and CABG has been assessed in many studies, with the vast majority reporting promising effects²⁸⁻³², some of which indicated a greater benefit for patients undergoing CABG^{29,33,34}. This is likely due to the more extensive surgical procedure with greater postoperative muscle deconditioning than with the less invasive PCI procedure, in which patients can ambulate immediately following the procedure. Therefore, CABG patients have a lower functional capacity at the entry of CR, but this phenomenon is reversible and transient with the aid of CR^{29,33}, emphasizing the importance of CR after CABG. In this study, the authors found that both groups of patients significantly improved after CR, although no significant difference was found between CABG and PCI in physical activity, left-ventricular EF, treadmill exercise test duration, and METs.

A study on PCI demonstrated that CR positively affected all aspects of the lipid profile level³⁵ with evidence that lipid profile components significantly decreased with CR following CABG³⁶. Although there is a lack of evidence for a link between exercise-based CR and fasting blood sugar (FBS) in patients with PCI³⁵, it was revealed that FBS and triglycerides (TG) decreased only in the PCI group with high-density lipoprotein cholesterol (HDL-C) increasing in both groups and no change in total cholesterol (TC) and low-density LDL-C with CR. Possible explanations are the worsening of insulin sensitivity by statins³⁵, patients' nutrition at home, their compliance with dietary recommendations, ethnic differences, the intensity of physical activity, and its duration. Besides, except for TG which was significantly decreased in patients with PCI, CR had the same effect on the lipid profile and FBS of CABG and PCI patients.

Both resting and peak heart rate (HR) significantly changed in both groups with no significantly greater change in favor of CABG or PCI patients. Other studies found a greater change in resting HR in patients with CABG than PCI, perhaps as an indicator of greater parasympathetic tone due to the longer convalescence period after surgery³⁷. Nevertheless, as HR-lowering drugs such as betablockers are prescribed to lower the heart demand Table 1. Cardiac rehabilitation participants' baseline characteristics before the program

Variables		Total (n=997)	PCI (n=440)	CABG (n=557)	Р
	Never	761 (76.32)	335 (76.13)	426 (76.48)	
Smoking	Current	93 (9.32)	45 (10.22)	48 (8.61)	0.61
n, (%)	Past	143 (14.34)	60 (13.63)	83 (14.9)	
Physical Activity	(MET.min/week)		00 (10100)	00 (1 lis)	
Walking		2025.87±2141.84	1885.80±2123.94	2212.63±2157.77	0.03
Moderate		1896.29±3832.13	2391.70±4670.37	1235.73±2118.79	< 0.0001
Vigorous		1140.09±4322.76	824.88±2551.24	1560.36±5894.96	0.86
Total		9265.11±5367.67	9451.47±5497.35	9016.25±5195.70	0.52
Lab Data					
Fasting Blood S		111.08 ± 36.63	112.08 ± 40.39	110.30±33.41	0.80
Triglyceride (m		165.54±91.49	163.87 ± 90.98	166.85±91.95	0.38
	oprotein (mg/dL)	92.85±35.57	86.66±32.26	97.78±37.30	< 0.0001
	ooprotein (mg/dL)	39.54±9.19	38.92 ± 8.58	40.03±9.62	0.14
Fotal cholester		167.03±45.85	160.33±44.43	172.31±46.29	< 0.0001
Cardiac Functio					
Ejection fractio		51.21±11.27	52.02±11.87	50.57±10.73	0.01
	Resting HR (bpm)	79.78±16.03	76.69±15.00	82.23±16.40	< 0.0001
	Peak HR (bpm)	125.87±23.85	122.52±23.19	128.53 ± 24.05	< 0.0001
	Resting SBP (mmHg)	116.99±17.66	117.07±16.48	116.94±18.57	0.83
Treadmill	Peak SBP (mmHg)	131.84±22.53	129.35±23.43	133.39 ± 21.85	0.01
Exercise	Resting DBP (mmHg)	72.25±10.24	72.72±9.68	71.86±10.67	0.20
stress	Peak DBP (mmHg)	77.43±10.52	76.84±10.98	77.79±10.23	0.40
test	Test Duration (min)	14.41±4.94	14.36±4.64	14.45 ± 5.18	0.67
	METs	8.49±3.06	8.79±3.32	8.25±2.81	0.01
	Negative	700 (70.21)	312 (70.9)	388 (69.65)	
	Result Positive	100 (10.03)	31 (7.07)	69 (12.38)	0.01 ^a
	UD	197 (19.75)	97 (22.04)	100 (17.95)	
Psychological st		511 (51.01)	200 ((5 45)	100 (55.04)	
Anxiety	Normal -mild	711 (71.31)	288 (65.45)	423 (75.94)	
Level	Moderate	231 (23.16)	123 (27.95)	108 (19.38)	< 0.0001
	Severe	51 (5.11)	26 (5.9)	25 (4.48)	
A mariata Sama	Very severe	4 (0.4)	3 (0.68)	1 (0.17)	<0.0001
Anxiety Score	Low	40.34±10.61	41.89±11.11	39.09±10.02	< 0.0001
	LUW	735 (73.72)	293 (66.59)	442 (79.35)	
Depression			65 (14 77)	52 (0 51)	
Depression	Mild	118 (11.83)	65 (14.77) 52 (11.81)	53 (9.51)	< 0.0001
Depression	Mild Intermediate	118 (11.83) 90 (9.02)	52 (11.81)	38 (6.82)	< 0.0001
level	Mild Intermediate High	118 (11.83) 90 (9.02) 54 (5.41)	52 (11.81) 30 (6.81)	38 (6.82) 24 (4.30)	
level Depression Scol	Mild Intermediate High re	118 (11.83) 90 (9.02) 54 (5.41) 11.19±9.14	52 (11.81) 30 (6.81) 12.50±10.06	38 (6.82) 24 (4.30) 10.13±8.18	<0.0001
Depression level Depression Scol	Mild Intermediate High re Physical functioning	118 (11.83) 90 (9.02) 54 (5.41) 11.19±9.14 59.05±23.33	52 (11.81) 30 (6.81) 12.50±10.06 60.78±24.79	38 (6.82) 24 (4.30) 10.13±8.18 56.96±21.28	<0.0001 0.02
level Depression Scor	Mild Intermediate High re Physical functioning Role-Health	118 (11.83) 90 (9.02) 54 (5.41) 11.19±9.14 59.05±23.33 35.31±37.25	52 (11.81) 30 (6.81) 12.50±10.06 60.78±24.79 36.92±37.98	38 (6.82) 24 (4.30) 10.13±8.18 56.96±21.28 33.36±36.31	<0.0001 0.02 0.26
Jepression Jevel Depression Scor General	Mild Intermediate High re Physical functioning Role-Health Body pain	$118 (11.83) 90 (9.02) 54 (5.41) 11.19\pm9.1459.05\pm23.3335.31\pm37.2563.47\pm26.24$	52 (11.81) 30 (6.81) 12.50±10.06 60.78±24.79 36.92±37.98 64.22±26.33	$\begin{array}{c} 38 \ (6.82) \\ 24 \ (4.30) \\ 10.13 \pm 8.18 \\ 56.96 \pm 21.28 \\ 33.36 \pm 36.31 \\ 62.56 \pm 26.14 \end{array}$	<0.0001 0.02 0.26 0.45
Jepression Jepression Scor General Quality of	Mild Intermediate High re Physical functioning Role-Health Body pain General health	$118 (11.83) 90 (9.02) 54 (5.41) 11.19\pm9.1459.05\pm23.3335.31\pm37.2563.47\pm26.2459.14\pm19.07$	$52 (11.81)$ $30 (6.81)$ 12.50 ± 10.06 60.78 ± 24.79 36.92 ± 37.98 64.22 ± 26.33 58.95 ± 18.96	$\begin{array}{c} 38\ (6.82)\\ 24\ (4.30)\\ 10.13\pm 8.18\\ 56.96\pm 21.28\\ 33.36\pm 36.31\\ 62.56\pm 26.14\\ 59.38\pm 19.22 \end{array}$	<0.0001 0.02 0.26 0.45 0.82
Depression Depression Scor General Quality of	Mild Intermediate High re Physical functioning Role-Health Body pain General health Energy/Fatigue	$118 (11.83) 90 (9.02) 54 (5.41) 11.19\pm9.1459.05\pm23.3335.31\pm37.2563.47\pm26.2459.14\pm19.0756.95\pm22.10$	$52 (11.81)$ $30 (6.81)$ 12.50 ± 10.06 60.78 ± 24.79 36.92 ± 37.98 64.22 ± 26.33 58.95 ± 18.96 56.45 ± 22.64	$\begin{array}{c} 38\ (6.82)\\ 24\ (4.30)\\ 10.13\pm 8.18\\ 56.96\pm 21.28\\ 33.36\pm 36.31\\ 62.56\pm 26.14\\ 59.38\pm 19.22\\ 57.57\pm 21.45 \end{array}$	<0.0001 0.02 0.26 0.45 0.82 0.42
Depression level Depression Scol	Mild Intermediate High re Physical functioning Role-Health Body pain General health Energy/Fatigue Social functioning	$118 (11.83) 90 (9.02) 54 (5.41) 11.19\pm9.1459.05\pm23.3335.31\pm37.2563.47\pm26.2459.14\pm19.0756.95\pm22.1067.46\pm25.93$	$52 (11.81)$ $30 (6.81)$ 12.50 ± 10.06 60.78 ± 24.79 36.92 ± 37.98 64.22 ± 26.33 58.95 ± 18.96 56.45 ± 22.64 68.50 ± 26.36	$\begin{array}{c} 38\ (6.82)\\ 24\ (4.30)\\ 10.13\pm 8.18\\ 56.96\pm 21.28\\ 33.36\pm 36.31\\ 62.56\pm 26.14\\ 59.38\pm 19.22\\ 57.57\pm 21.45\\ 66.19\pm 25.39 \end{array}$	<0.0001 0.02 0.26 0.45 0.82 0.42 0.18
Depression Depression Scor General Quality of	Mild Intermediate High re Physical functioning Role-Health Body pain General health Energy/Fatigue Social functioning Role emotional	$118 (11.83) 90 (9.02) 54 (5.41) 11.19\pm9.1459.05\pm23.3335.31\pm37.2563.47\pm26.2459.14\pm19.0756.95\pm22.1067.46\pm25.9353.75\pm40.93$	$52 (11.81)$ $30 (6.81)$ 12.50 ± 10.06 60.78 ± 24.79 36.92 ± 37.98 64.22 ± 26.33 58.95 ± 18.96 56.45 ± 22.64 68.50 ± 26.36 54.99 ± 41.61	$\begin{array}{c} 38 \ (6.82) \\ 24 \ (4.30) \\ 10.13 \pm 8.18 \\ 56.96 \pm 21.28 \\ 33.36 \pm 36.31 \\ 62.56 \pm 26.14 \\ 59.38 \pm 19.22 \\ 57.57 \pm 21.45 \\ 66.19 \pm 25.39 \\ 52.24 \pm 40.10 \end{array}$	<0.0001 0.02 0.26 0.45 0.82 0.42 0.18 0.37
Depression Depression Scor General Quality of life	Mild Intermediate High re Physical functioning Role-Health Body pain General health Energy/Fatigue Social functioning Role emotional Emotional Well being	$118 (11.83) 90 (9.02) 54 (5.41) 11.19\pm9.1459.05\pm23.3335.31\pm37.2563.47\pm26.2459.14\pm19.0756.95\pm22.1067.46\pm25.9353.75\pm40.9366.11\pm22.30$	$52 (11.81)$ $30 (6.81)$ 12.50 ± 10.06 60.78 ± 24.79 36.92 ± 37.98 64.22 ± 26.33 58.95 ± 18.96 56.45 ± 22.64 68.50 ± 26.36 54.99 ± 41.61 66.57 ± 21.79	$\begin{array}{c} 38 \ (6.82) \\ 24 \ (4.30) \\ 10.13\pm 8.18 \\ 56.96\pm 21.28 \\ 33.36\pm 36.31 \\ 62.56\pm 26.14 \\ 59.38\pm 19.22 \\ 57.57\pm 21.45 \\ 66.19\pm 25.39 \\ 52.24\pm 40.10 \\ 65.55\pm 22.93 \end{array}$	<0.0001 0.02 0.26 0.45 0.82 0.42 0.18 0.37 0.64
Depression level Depression Scor General Quality of life Health-	Mild Intermediate High re Physical functioning Role-Health Body pain General health Energy/Fatigue Social functioning Role emotional Emotional Well being Physical	$118 (11.83) 90 (9.02) 54 (5.41) 11.19\pm9.1459.05\pm23.3335.31\pm37.2563.47\pm26.2459.14\pm19.0756.95\pm22.1067.46\pm25.9353.75\pm40.9366.11\pm22.304.75\pm1.08$	$52 (11.81)$ $30 (6.81)$ 12.50 ± 10.06 60.78 ± 24.79 36.92 ± 37.98 64.22 ± 26.33 58.95 ± 18.96 56.45 ± 22.64 68.50 ± 26.36 54.99 ± 41.61 66.57 ± 21.79 4.69 ± 1.10	$\begin{array}{c} 38 \ (6.82) \\ 24 \ (4.30) \\ 10.13\pm 8.18 \\ 56.96\pm 21.28 \\ 33.36\pm 36.31 \\ 62.56\pm 26.14 \\ 59.38\pm 19.22 \\ 57.57\pm 21.45 \\ 66.19\pm 25.39 \\ 52.24\pm 40.10 \\ 65.55\pm 22.93 \\ 4.79\pm 1.07 \end{array}$	<0.0001 0.02 0.26 0.45 0.82 0.42 0.18 0.37 0.64 0.28
Depression level Depression Scou General Quality of	Mild Intermediate High re Physical functioning Role-Health Body pain General health Energy/Fatigue Social functioning Role emotional Emotional Well being	$118 (11.83) 90 (9.02) 54 (5.41) 11.19\pm9.1459.05\pm23.3335.31\pm37.2563.47\pm26.2459.14\pm19.0756.95\pm22.1067.46\pm25.9353.75\pm40.9366.11\pm22.30$	$52 (11.81)$ $30 (6.81)$ 12.50 ± 10.06 60.78 ± 24.79 36.92 ± 37.98 64.22 ± 26.33 58.95 ± 18.96 56.45 ± 22.64 68.50 ± 26.36 54.99 ± 41.61 66.57 ± 21.79	$\begin{array}{c} 38 \ (6.82) \\ 24 \ (4.30) \\ 10.13\pm 8.18 \\ 56.96\pm 21.28 \\ 33.36\pm 36.31 \\ 62.56\pm 26.14 \\ 59.38\pm 19.22 \\ 57.57\pm 21.45 \\ 66.19\pm 25.39 \\ 52.24\pm 40.10 \\ 65.55\pm 22.93 \end{array}$	0.26 0.45 0.82 0.42 0.18 0.37 0.64

PCI, percutaneous coronary intervention; CABG, coronary artery bypass grafting surgery; UD, undetermined; HR, heart rate;SBP, systolic blood pressure; DBP, diastolic blood pressure

^a According to Bonferroni method analysis, this significant P value was seen in two situation: when comparing negative group with positive group and when comparing group "other" with positive group. ^b According to Bonferroni method analysis, this significant P value was seen when comparing the normal-mild group with

moderate group.

^c According to Bonferroni method analysis, this significant P value was seen when comparing group "low" with other groups.

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Table 2.	Comp	arison of	variables	before	and after	the pro	gram in e	ach group
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Variables		PCI Before	After	Р	CABG Before	After	Р
Smoking	Never Current Past	335 (76.13) 45 (10.22) 60 (13.63)	335 (76.13) 12 (2.72) 93 (21.13)	<0.0001ª	426 (76.48) 48 (8.61) 83 (14.9)	426 (76.48) 16 (2.87) 115 (20.64)	<0.0001
	<i>ivity</i> (MET.min/w			0.0004			0.0004
Walking		1885.80±2123.94	2273.18±1956.64	< 0.0001	2212.63±2157.77	3094.22±2797.82	< 0.0001
Moderate		2391.70±4670.37	3399.20±3059.22	<0.0001 <0.0001	1235.73±2118.79 1560.36±5894.96	3080.00±3686.04	<0.0001 0.16
Vigorous Total		824.88±2551.24 9451.47±5497.35	2346.86±9522.90 11179.09±5078.57	< 0.0001	9016.25±5195.70	1293.04±4097.96 11218.78±5275.16	< 0.0001
Lad Data		J+J1.+/±J+//.55	11177.09±3078.37	<0.0001	9010.25±5195.70	11210.76±5275.10	<0.0001
Fasting Bloo (mg/dL)	od Sugar	112.08±40.39	107.31±30.56	0.01	110.30±33.41	108.97±33.44	0.84
Triglyceride	e (mg/dL)	163.87±90.98	144.94±65.72	0.01	166.85±91.95	156.97±76.48	0.66
	lipoprotein	86.66±32.26	84.48±27.30	0.81	97.78±37.30	94.38±31.69	0.37
(mg/dL)		00.00±32.20	04.40±27.50	0.01	J1.10±31.30)4.50±51.07	0.57
	y lipoprotein	38.92±8.58	39.89±10.46	0.03	40.03±9.62	41.41±9.77	0.01
(mg/dL) Total abolas	terol (mg/dL)	160.33±44.43	154.68±36.03	0.26	172.31±46.29	167.50±37.84	0.44
Cardiac Fun		100.35±44.45	134.08±30.03	0.20	1/2.31±40.29	107.30±37.84	0.44
Ejection fra		52.02±11.87	53.79±10.51	< 0.0001	50.57±10.73	53.96±9.59	< 0.000
ejection fra	Resting HR						
	(bpm)	76.69±15.00	74.47±14.42	0.02	82.23±16.40	77.13±15.84	< 0.000
	Peak HR	122.52±23.19	131.36±23.68	< 0.0001	128.53±24.05	130.03±24.12	0.01
	(bpm)	122.32±23.19	131.30±23.08	<0.0001	128.35±24.05	130.03±24.12	0.01
	Resting SBP (mmHg)	117.07±16.48	113.70±16.12	0.02	116.94±18.57	116.05±17.02	0.04
	Peak SBP (mmHg)	129.35±23.43	129.35±21.49	0.85	133.39±21.85	134.68±25.45	0.53
Freadmill Exercise	Resting DBP (mmHg)	72.72±9.68	71.42±9.05	0.48	71.86±10.67	72.11±9.97	0.20
stress test	Peak DBP (mmHg)	76.84±10.98	76.82±10.41	0.47	77.79±10.23	78.42±15.13	0.13
	Test Duration	14.36±4.64	18.09±4.95	< 0.0001	14.45±5.18	17.73±4.76	<0.000
	(min) METs	8.79±3.32	11.93±3.70	< 0.0001	8.25±2.81	10.90±3.07	< 0.000
	Negative	312 (70.9)	379 (86.13)	<0.0001	388 (69.65)	474 (85.09)	<0.000
	Result Positive	31 (7.07)	16 (3.63)	<0.0001 ^b	69 (12.38)	20 (3.59)	< 0.000
	UD	97 (22.04)	45 (10.22)	-0.0001	100 (17.95)	63 (11.31)	-0.000
Psychologica		97 (22.04)	45 (10.22)		100 (17.95)	05 (11.51)	
esychologici	Normal -mild	288 (65.45)	342 (77.72)		423 (75.94)	459 (82.4)	
Anxiety	Moderate	123 (27.95)	83 (18.86)		108 (19.38)	93 (16.69)	
Level	Severe	26 (5.9)	15 (3.4)	<0.0001°	25 (4.48)	5 (0.89)	< 0.000
	Very severe	3 (0.68)	0(0)		1 (0.17)	0 (0)	
Anxiety Sco		41.89±11.11	39.72±11.13	< 0.0001	39.09±10.02	38.02±9.78	0.000
•	Low	293 (66.59)	357 (81.13)		442 (79.35)	496 (89.04)	
Depression	Mild	65 (14.77)	43 (9.77)	in coord	53 (9.51)	27 (4.84)	-0.000
level	Intermediate	52 (11.81)	29 (6.59)	<0.0001 ^d	38 (6.82)	25 (4.48)	< 0.000
	High	30 (6.81)	11 (2.5)		24 (4.30)	9 (1.61)	
Depression S	Score	12.50±10.06	10.31±9.16	< 0.0001	10.13±8.18	8.18±7.34	< 0.000
	Physical functioning	60.78±24.79	70.16±20.86	< 0.0001	56.96±21.28	68.13±20.93	<0.000
General	Role-Health	36.92 ± 37.98	55.43±39.35	< 0.0001	33.36±36.31	51.03 ± 38.60	< 0.000
eneral Quality of	Body pain	64.22±26.33	74.39 ± 22.80	< 0.0001	62.56±26.14	73.58±21.67	< 0.000
fe	General health	58.95 ± 18.96	64.03±18.74	< 0.0001	59.38±19.22	64.38±17.82	< 0.000
	Energy/Fatigue	56.45±22.64	64.16±20.19	< 0.0001	57.57±21.45	62.65±20.56	< 0.000
	Social	68.50±26.36	77.60±22.02	< 0.0001	66.19±25.39	76.21±22.63	< 0.000
	functioning			0.0001			2.000

** * * * *		PCI			CABG		
Variables		Before	After	Р	Before	After	Р
	Role emotional	54.99±41.61	64.86±38.30	< 0.0001	52.24±40.10	64.66±37.56	< 0.0001
	Emotional Well-being	66.57±21.79	71.77±19.56	< 0.0001	65.55±22.93	69.74±21.04	< 0.0001
Health	Physical	4.69±1.10	5.25±1.00	< 0.0001	4.79±1.07	5.27±0.93	< 0.0001
related	Social	4.78±1.06	5.35±1.01	< 0.0001	4.84±1.09	5.38 ± 0.96	< 0.0001
Quality of life	Emotional	4.64±0.86	4.88±0.80	< 0.0001	4.83±0.98	5.07±0.86	< 0.0001

Continued Table 2. Comparison of variables before and after the program in each group

PCI, percutaneous coronary intervention; CABG, coronary artery bypass grafting surgery; UD, undetermined; HR, heart rate; SBP, systolic blood pressure; DBP, diastolic blood pressure

^a Bonferroni correction showed significant difference when comparing the "never" group with either "smoker" or "past" group.

^b Bonferroni correction showed significant difference when comparing the "negative" group with either "positive" or "other" group. ^c Bonferroni correction showed significant difference when comparing the "normal-mild" group with either "moderate" or "severe" group.

group. ^d Bonferroni correction showed significant difference when comparing the "low" group with either "mild", "intermediate" or, "high" group.

	Table 3. Com	parison of	delta difference	e of each	variable	between the groups
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			PCI	CABG	
Variables		Total	[After-Before]	[After-Before]	Р*
Quit Smoking n, (65 (6.5)	33 (7.5)	32 (5.7)	0.10
Physical Activity (N	/IET.min/week)				
Walking		536.37±2672.43	287.84±2533.96	834.62±2810.94	0.15 ^a
Moderate		1421.28±4710.96	1054.29±5385.52	1861.66 ± 3718.88	0.86
Vigorous		964.85±7651.86	1533.70±9422.53	282.24±4672.63	0.26
Total		2117.79±6696.30	1995.71±6610.77	2262.70±6820.68	0.58
Lab Data					
Fasting Blood Sug	ar (mg/dL)	-2.66 ± 28.48	-4.81 ± 28.64	-0.88 ± 28.26	0.31
Triglyceride (mg/d		-7.96 ± 74.78	-11.76 ± 61.54	-4.81 ± 84.12	0.01 ^a
Low-density lipop	rotein (mg/dL)	-1.26 ± 30.88	-0.47 ± 27.43	-1.92 ± 33.54	0.37
High-density lipop	rotein (mg/dL)	1.01±9.11	1.01 ± 8.13	1.01 ± 9.86	0.85
Total cholesterol (-2.57±38.10	-3.48 ± 37.13	-1.82 ± 38.92	0.11
Cardiac Function	'ests				
Ejection fraction (%)	2.51±6.74	2.03±6.47	2.93±6.94	0.14
	Resting HR (bpm)	-3.31±14.42	-1.95 ± 14.87	-4.46±13.95	0.12
	Peak HR (bpm)	5.83±23.83	8.96±25.74	3.15±21.75	0.06
	Resting SBP (mmHg)	-2.30 ± 17.96	-2.60 ± 16.60	-2.04 ± 19.04	0.31
Treadmill	Peak SBP (mmHg)	1.03 ± 22.37	0.01±21.97	1.70 ± 22.64	0.81
Exercise	Resting DBP (mmHg)	-0.70 ± 10.73	-0.66 ± 10.71	-0.73±10.77	0.27
stress	Peak DBP (mmHg)	1.30±14.31	1.42±13.02	1.22±15.12	0.34
test	Test Duration (min)	3.52±4.41	3.53±4.28	3.51±4.53	0.27
	METs	2.73±2.64	2.98±2.89	2.53±2.39	0.98
	Get Negative result n,	153 (15.34)	67 (15.22)	86 (15.43)	0.72
	(%)	· · ·	~ /	× /	
Psychological statu	15	1.00+0.21	2 24+0 11	1 70 17 57	0.57
Anxiety Score		-1.99 ± 8.31	-2.24±9.11	-1.78±7.57	0.37
Depression Score	Dhamia al fam ati anima	-2.04±6.54	-1.71±6.63	-2.30±6.45	
	Physical functioning	10.07±22.62	10.58±22.65	9.49±22.64	0.60
	Role-Health	18.74±43.96	20.23±44.55	17.03±43.32	0.22
General	Body pain General health	9.90±25.56	9.59±24.95	10.26±26.29	0.09 0.56
		3.96±17.88	4.00±17.02	3.91±18.85	
Quality of Life	Energy/Fatigue	5.71±19.35	7.30±18.07	3.89±20.59	0.19
	Social functioning	8.13±24.29	8.67±22.25	7.52±26.45	0.40^{a}
	Role emotional	11.45±47.55	11.93±45.95	10.91±49.42	0.97
	Emotional Well being	3.82±18.34	4.42±17.27	3.14±19.51	0.63
Health related	Physical	0.46±0.86	0.48±0.86	0.45±0.86	0.58
Quality of Life	Social	0.51±0.90	0.52±0.8	0.51±0.92	0.42
	Emotional	0.22±0.76	0.22±0.74	0.22±0.78	0.33ª

PCI, percutaneous coronary intervention; CABG, coronary artery bypass grafting surgery; HR, heart rate; SBP, systolic blood pressure; DBP, diastolic blood pressure

*All p-values were obtained by ANCOVA, except for quit smoking obtained by Logistic regression.

^a Based on logarithmic transformation due to heterogeneity of variance.

after any coronary events, therefore, HR change will be under drug control rather than a CR response.

Resting systolic blood pressure (SBP) decreased significantly and equally in both groups, however, diastolic blood pressure (DBP) and peak SBP were not affected by CR. Although some studies support the authors' findings that exercise-based CR does not influence blood pressure (BP) in patients with either PCI or CABG,^{37,38} it was suggested that CABG patients had significantly lower peak DBP as well as resting and peak SBP in comparison with the group without CR³⁹ and CR participants after PCI had significantly lower SBP and DBP³⁵. These hemodynamic contradictions can be due to different exercise protocols with various intensities, age and gender differences, sample size variations, and medications after each procedure³⁹.

Improvement in controlling anxiety and depression, along with enhanced general quality of life (QoL) and health-related quality of life (HR-QoL), are among the established outcomes of the CR program⁴⁰⁻⁴² and were observed in the present study. However, no significant difference was found between the two intervention groups. Additionally, it has been shown that patients who underwent PCI have better HR-QoL in the short-term following CR than those who underwent CABG43. Furthermore, it was suggested that, in contrast to the CABG patients, PCI patients would have better HR-QoL after the intervention and before the CR, suggesting that greater improvement may be observed in CABG than in PCI44. Although it remains controversial, these findings are linked to possible confounding factors like age, sex, socioeconomic status, education level, body weight, and comorbid disease^{43,45}.

This study could have been limited by the fact that medical documents of one CR referral center were reviewed, and socioeconomic status, educational level, and logistic factors were not evaluated. According to the authors' observation, although CR is advised after both PCI and CABG, more CABG patients participated due to the low PCI referral rate. Moreover, the retrospective nature of the study should be taken into account.

Conclusion

Both PCI and CABG patients from the Eastern Mediterranean region benefit significantly, and to the same extent, from CR. Therefore, it indicates that CR should be supported by the healthcare insurances, noticed by policymakers, and recommended by the physician to both groups.

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Conflict of Interest

All authors declare no potential conflict of interest.

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