

Clinical Trial Protocol

Iranian Registry of Clinical Trials

07 Jun 2026

Investigating the effect of different exercise regimens of the first phase of cardiac rehabilitation on the discharge day performance of open heart surgery patients

Protocol summary

Study aim

Investigating the effect of different exercise regimens of the first phase of cardiac rehabilitation on the discharge day performance of open heart surgery patients

Design

Clinical trial with a control group, with parallel groups, single-blind, randomized, on 32 patients. Simple randomization was used.

Settings and conduct

This study will be conducted in Erfan Niayesh Hospital. First, teaching patients about the effects of surgery on cardiopulmonary function, general cases after surgery one day before surgery and the cardiac rehabilitation program, which generally includes mobilization of the patient as soon as possible, teaching breathing exercises, personal care will be given from the day before the surgery to discharge. The patient and the therapist know the study groups and only the evaluator (examiner) is blind. Before surgery and discharge day, spirometry and 6-minute walking test were taken by an evaluator.

Participants/Inclusion and exclusion criteria

People who candidates for open heart surgery (coronary artery bypass graft, valve replacement, or both) based on: - Age over 18 years and less than 70 years - Independent walking - No history of surgery in the past - Non-emergency surgery - No history of recent myocardial infarction (past 6 months) - Not having a pacemaker - Not having chronic obstructive pulmonary disease - Absence of severe non-cardiac disease

Intervention groups

The cardiac rehabilitation program, which generally includes: mobilization of the patient as soon as possible, teaching breathing exercises and personal care, once or twice daily (one group, once and the other group twice a day), from the day before the operation until the day of discharge will be given.

Main outcome variables

distance walked in six minute walk tes: forced vital capacity :Forced expiratory volume in first second:forced expiratory volume in first second/forced vital capacity ratio

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20230527058305N1**
Registration date: **2023-06-21, 1402/03/31**
Registration timing: **prospective**

Last update: **2023-06-21, 1402/03/31**

Update count: **0**

Registration date

2023-06-21, 1402/03/31

Registrant information

Name

Oweis Rezaei

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 21 4979 6260

Email address

oweisrezaei@gmail.com

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2023-06-25, 1402/04/04

Expected recruitment end date

2023-07-28, 1402/05/06

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Investigating the effect of different exercise regimens of the first phase of cardiac rehabilitation on the discharge day performance of open heart surgery patients

Public title

Investigating the effect of different exercise regimens of the first phase of cardiac rehabilitation

Purpose

Treatment

Inclusion/Exclusion criteria**Inclusion criteria:**

Open heart surgery(coronary artery bypass graft, valve replacement or both) candidate patients

Exclusion criteria:

History of open heart surgery Emergency surgery History of recent myocardial infarction (past 6 months) Implanted pace maker Having chronic obstructive pulmonary disease Severe non-cardiac disease

Age

From **18 years** old to **70 years** old

Gender

Both

Phase

N/A

Groups that have been masked

- Outcome assessor

Sample size

Target sample size: **32**

Randomization (investigator's opinion)

Randomized

Randomization description

A simple randomization method was used to select people in these two groups. From the beginning, it was agreed that intervention group 1 would be given odd numbers and intervention group 2 would be given even numbers. Then, according to the number of sample size (32) under study, the relevant random numbers were extracted using Excel software, and each number was written on a card and placed in an envelope, and the envelopes were sealed, and on each envelope was the patient's number. Envelope number 1 will be given to the first patient who is enrolled in the study, envelope number 2 will be given to second patient, and so on until the end of the study sample.

Blinding (investigator's opinion)

Single blinded

Blinding description

In this study patients and therapist know the situation of patients in groups. Only assessor do not know the situation of patients in groups.

Placebo

Not used

Assignment

Parallel

Other design features**Secondary Ids**

empty

Ethics committees**1****Ethics committee****Name of ethics committee**

Ethics committee of University of Social Welfare and Rehabilitation sciences

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Kodakyaar Deadend., Daneshjoo Blvd., Evin., Tehran

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1985713834

Approval date

2023-06-14, 1402/03/24

Ethics committee reference number

IR.USWR.REC.1402.047

Health conditions studied**1****Description of health condition studied**

Chronic ischemic heart disease

ICD-10 code

I25

ICD-10 code description

Chronic ischemic heart disease

2**Description of health condition studied**

Rheumatic mitral valve diseases

ICD-10 code

I05

ICD-10 code description

Rheumatic mitral valve diseases

3**Description of health condition studied**

Nonrheumatic mitral valve disorders

ICD-10 code

I34

ICD-10 code description

Nonrheumatic mitral valve disorders

4**Description of health condition studied**

Multiple valve diseases

ICD-10 code

I08

ICD-10 code description

Multiple valve diseases

5

Description of health condition studied

Rheumatic aortic valve diseases

ICD-10 code

I06

ICD-10 code description

Rheumatic aortic valve diseases

6

Description of health condition studied

Nonrheumatic aortic valve disorders

ICD-10 code

I35

ICD-10 code description

Nonrheumatic aortic valve disorders

Primary outcomes

1

Description

distance walk in six minute walk test

Timepoint

Before intervention (surgery) and discharge day

Method of measurement

In this test we want from patient to walk as far as he or she can in six minute . Patient will walk in a 30 meter path up and down. There are some chairs through the path for rest.patient allowed to rest in case of fatigue or symptoms like chest pain, shortness of breath or musculoskeletal pain.the patient continues to walk until the time of six minutes is over.

2

Description

Forced vital capacity

Timepoint

Before intervention (surgery) and discharge day

Method of measurement

With spirometer (spirolab MIR company)

3

Description

Forced expiratory volume in first second

Timepoint

Before intervention (surgery) and discharge day

Method of measurement

With spirometer (spirolab MIR company)

Secondary outcomes

empty

Intervention groups

1

Description

Intervention group:cardiac rehabilitation program, which generally includes: early mobilization of the patient as soon as possible, teaching breathing exercises and organs, teaching personal care, will be given in the number of one session per day, from the day before the operation to the day of discharge.

Category

Rehabilitation

2

Description

Intervention group:cardiac rehabilitation program, which generally includes: early mobilization of the patient as soon as possible, teaching breathing exercises and organs, teaching personal care, will be given in the number of two session per day, from the day before the operation to the day of discharge.(The difference between this group and intervention group 1 is in the frequency of therapeutic sessions, which is twice a day).

Category

Rehabilitation

Recruitment centers

1

Recruitment center

Name of recruitment center

Erfan niayesh hospital

Full name of responsible person

Oweis Rezaei

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

University of social welfare and rehabilitation sciences

Full name of responsible person

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Grant name
Grant code / Reference number
Is the source of funding the same sponsor organization/entity?
Yes
Title of funding source
University of social welfare and rehabilitation sciences
Proportion provided by this source
100
Public or private sector
Public
Domestic or foreign origin
Domestic
Category of foreign source of funding
empty
Country of origin
Type of organization providing the funding
Academic

Person responsible for general inquiries

Contact
Name of organization / entity
University of social welfare and rehabilitation sciences
Full name of responsible person
Oweis Rezaei
Position
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Sharing plan

Deidentified Individual Participant Data Set (IPD)
Yes - There is a plan to make this available
Study Protocol
Undecided - It is not yet known if there will be a plan to make this available
Statistical Analysis Plan
Undecided - It is not yet known if there will be a plan to make this available
Informed Consent Form
No - There is not a plan to make this available

Clinical Study Report

Undecided - It is not yet known if there will be a plan to make this available

Analytic Code

Undecided - It is not yet known if there will be a plan to make this available

Data Dictionary

Undecided - It is not yet known if there will be a plan to make this available

Title and more details about the data/document

All the data related to the demographic information of the patients and their related medical documents and the test results of this research can be shared.

When the data will become available and for how long

The access period starts after the article is published in the journal

To whom data/document is available

Researchers working in academic and scientific institutions

Under which criteria data/document could be used

Research uses for articles in the medical field

From where data/document is obtainable

1. Oweis Rezaei at the email address oweisrezaei@gmail.com at the phone number 0098 939 374 6002 2. Dr. Farhad Azadi at the email address fa_azadi@yahoo.com

What processes are involved for a request to access data/document

First, the request is sent by the applicant to the email address of the mentioned people. Then, if the main researcher approves and after reviewing the applicant's documents, the requested data will be sent as a file

Comments