

Clinical Trial Protocol

Iranian Registry of Clinical Trials

28 Jun 2026

Effect of education through group discussion on self-care behaviors promotion in patients with heart failure

Protocol summary

Summary

This will be a randomized clinical trial. The aim of the study will determine the effect of group discussion on Self-care behaviors promotion in patients with heart failure. Introduction Patients of CHF diagnosed will be included. Patients with hospital readmissions during the study, Unwillingness to continue participating in the study, no participation in one training session will be excluded. The sample size of 80 will be required. Before education the questionnaire will be given to the control and experience groups. Education program will be set group discussion by the investigator. Samples will come to the hospitals after 3 months of training and Questionnaire will be given to them again. Comparison of pre- and post-intervention difference between the means as the effect of group discussion on Self-care behaviors promotion in patients with heart failure will be considered in experimental and control groups.

General information

Acronym

Education through group discussion and heart failure

IRCT registration information

IRCT registration number: **IRCT2015120618464N3**

Registration date: **2016-02-16, 1394/11/27**

Registration timing: **prospective**

Last update:

Update count: **0**

Registration date

2016-02-16, 1394/11/27

Registrant information

Name

Soghra Nikpour

Name of organization / entity

Iran University of Medical Sciences

Country

Iran (Islamic Republic of)

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Email address

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Recruitment status

Recruitment complete

Funding source

Deputy of Research Iran University of Medical Sciences

Expected recruitment start date

2016-04-04, 1395/01/16

Expected recruitment end date

2016-08-06, 1395/05/16

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Effect of education through group discussion on self-care behaviors promotion in patients with heart failure

Public title

Effect of education through group discussion on self-care behaviors promotion in patients with heart failure to educational hospitals of Iran University of Medical Sciences.

Purpose

Prevention

Inclusion/Exclusion criteria

Inclusion criteria: The diagnosis of heart failure by doctor:age more than 18:A history of heart failure at least 6 months prior to study:Having at least read and writes in Persian: having an active phone line Exclusion criteria: Hospital readmission samples at the time of investigation:No participation in one training sessions:Unwillingness to continue participating in the

study

Age

No age limit

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: 80

Randomization (investigator's opinion)

Randomized

Randomization description

Blinding (investigator's opinion)

Single blinded

Blinding description

Placebo

Not used

Assignment

Parallel

Other design features

Eighty patients with heart failure randomly will be assigned and by random blocks will be placed in two experience and control groups.

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethics committee of University of Medical Sciences

Street address

Iran University of Medical Sciences, Shahid Hemmat Highway

City

Tehran

Postal code

1449614535

Approval date

2015-05-20, 1394/02/30

Ethics committee reference number

IR.IUMS.REC.1394.25732

Health conditions studied

1

Description of health condition studied

Congestive heart failure

ICD-10 code

I50.0

ICD-10 code description

Right ventricular failure (secondary to left heart failure)

2

Description of health condition studied

Left ventricular failure

ICD-10 code

I50.1

ICD-10 code description

Cardiac asthma ,Left heart failure ,Oedema of lung, Pulmonary oedemawith mention of heart disease NOS or heart failure

3

Description of health condition studied

Heart failure, unspecified

ICD-10 code

I50.9

ICD-10 code description

Cardiac, heart or myocardial failure NOS

Primary outcomes

1

Description

Self-care behaviors promotion

Timepoint

Before and 3 months after education

Method of measurement

Questionnaire

Secondary outcomes

empty

Intervention groups

1

Description

The experience group is given questionnaire before the intervention. They will be training three sessions for 90 minutes. After the intervention, 3 months later the will be given re-questionnaire.

Category

Prevention

2

Description

The control group is given questionnaire before the intervention; there is no intervention in the control group. After the intervention, 3 months later the control group will be given re-questionnaire.

Category

Prevention

Recruitment centers

1

Recruitment center

Name of recruitment center

Rajaei Cardiovascular, Medical & Research Center

Full name of responsible person

Dr Fereydoon Nouhi

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Web page address**Person responsible for scientific inquiries****Contact****Name of organization / entity**

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Full name of responsible person

Soghra Nikpour

Position

Assistant professor

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Web page address**Sponsors / Funding sources****1****Sponsor****Name of organization / entity**

Vice Chancellor for research of Iran University of Medical Sciences

Full name of responsible person

Dr Seyed Ali Javad Moosavi

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Iran University of Medical Sciences, Hemmat Highway

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Tehran

Grant name**Grant code / Reference number****Is the source of funding the same sponsor organization/entity?**

Yes

Title of funding source

Vice Chancellor for research of Iran University of Medical Sciences

Proportion provided by this source

100

Public or private sector*empty***Domestic or foreign origin***empty***Category of foreign source of funding***empty***Country of origin****Type of organization providing the funding***empty***Person responsible for general inquiries****Contact****Name of organization / entity**

Iran University of Medical Sciences

Full name of responsible person

Soghra Nikpour

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Person responsible for updating data**Contact****Name of organization / entity**

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Full name of responsible person

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Web page address**Sharing plan****Deidentified Individual Participant Data Set (IPD)***empty***Study Protocol**

empty
Statistical Analysis Plan
empty
Informed Consent Form
empty
Clinical Study Report

empty
Analytic Code
empty
Data Dictionary
empty