

Clinical Trial Protocol

Iranian Registry of Clinical Trials

04 Jun 2026

Study of the effect of spiritual intervention on quality of life and life expectancy of women after hysterectomy

Protocol summary

Study aim

Determining the effect of spiritual intervention on quality of life and life expectancy of women after hysterectomy

Design

This study is non-blinded clinical trial with control group. The study population will be included all women candidates for hysterectomy referring to women's ward of Imam Reza hospital of Kermanshah. 60 eligible patients will be selected conveniently and randomly will be assigned to intervention and control groups.

Settings and conduct

This study which will be conducted in Imam Reza hospital of Kermanshah is one-blinded one. Before the study and 4 weeks after the last session, the questionnaire of quality of life and life expectancy will be provided to patients, and both groups will be compared.

Participants/Inclusion and exclusion criteria

Inclusion criteria: Literacy; Not participate in similar classes; Do not have mental illness; Not being treated with chemotherapy and radiotherapy Exclusion criteria: Do not attend more than two sessions in intervention; Experience a psychological crisis during the study

Intervention groups

The intervention group for 8 sessions of 1.5 hours (weekly sessions) will receive group spiritual intervention with groups of 12 to 15 people. The control group will not receive any intervention

Main outcome variables

Quality of life; Life expectancy

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20181223042078N1**

Registration date: **2019-06-03, 1398/03/13**

Registration timing: **retrospective**

Last update: **2019-06-03, 1398/03/13**

Update count: **0**

Registration date

2019-06-03, 1398/03/13

Registrant information

Name

Mahin Hemati

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 83 3729 2820

Email address

mahinhemati1367@gmail.com

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2019-02-18, 1397/11/29

Expected recruitment end date

2019-03-20, 1397/12/29

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Study of the effect of spiritual intervention on quality of life and life expectancy of women after hysterectomy

Public title

Study of the effect of spiritual intervention on quality of life and life expectancy of women after hysterectomy

Purpose

Supportive

Inclusion/Exclusion criteria

Inclusion criteria:

Literacy Not participate in similar classes Not have a psychological crisis during the study Not treated with chemotherapy and radiotherapy Do not use drugs Do not have mental illness

Exclusion criteria:

Do not attend more than two sessions in intervention Experience a psychological crisis during the study

Age

From **20 years** old to **70 years** old

Gender

Female

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **60**

Randomization (investigator's opinion)

Randomized

Randomization description

Simple accident

Blinding (investigator's opinion)

Not blinded

Blinding description

Placebo

Not used

Assignment

Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethics committee of Kermanshah University of Medical Sciences

Street address

Vice Chancellor for Research Affairs, Kermanshah University of Medical Sciences, Building No.2, Shahid Beheshti

City

kermanshah

Province

Kermanshah

Postal code

6715847141

Approval date

2018-12-05, 1397/09/14

Ethics committee reference number

IR.KUMS.REC.1397.702

Health conditions studied

1

Description of health condition studied

Hysterectomy

ICD-10 code

Z90.7

ICD-10 code description

Acquired absence of genital organ(s)

Primary outcomes

1

Description

Quality of Life

Timepoint

Before the intervention begins, 2 months after the intervention begins

Method of measurement

Based on World Health Organization Quality of Life Questionnaire

2

Description

Hope of Life

Timepoint

Before the intervention begins, 2 months after the intervention begins

Method of measurement

Base on Snyder's Hope Scale

Secondary outcomes

empty

Intervention groups

1

Description

The intervention group for 8 sessions of 1.5 hours (weekly sessions) will received group spiritual intervention with groups of 12 to 15 people.

Category

Lifestyle

2

Description

The control group will not receive any intervention

Category

Other

Recruitment centers

1

Recruitment center

Name of recruitment center

Emam Reza Hospital

Full name of responsible person

Mahin Hemmati
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Emam Reza Hospital, Parastar Boulevard
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Student of Psychiatric Nursing
Latest degree
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Sponsors / Funding sources

1

Sponsor

Name of organization / entity
Kermanshah University of Medical Sciences
Full name of responsible person
Dr. Amir Jalali
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Government of the Faculty of Nursing and Midwifery
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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
Kermanshah University of Medical 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
Kermanshah University of Medical Sciences
Full name of responsible person
Mahin Hemmati
Position

Person responsible for scientific inquiries

Contact

Name of organization / entity
Kermanshah University of Medical Sciences
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Dr. Amir Jalali
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Person responsible for updating data

Contact

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Sharing plan**Deidentified Individual Participant Data Set (IPD)**

No - There is not a plan to make this available

Justification/reason for indecision/not sharing IPD

No more information.

Study Protocol

No - There is not a plan to make this available

Statistical Analysis Plan

No - There is not a plan to make this available

Informed Consent Form

No - There is not a plan to make this available

Clinical Study Report

No - There is not a plan to make this available

Analytic Code

No - There is not a plan to make this available

Data Dictionary

No - There is not a plan to make this available