

Clinical Trial Protocol

Iranian Registry of Clinical Trials

27 Jun 2026

Comparison the effectiveness of group mindfulness-based stress reduction and group cognitive-behavioral stress management on biological markers and psychological symptoms in patients with essential hypertension

Protocol summary

Summary

The purpose of this study is comparison the effectiveness of group mindfulness-based stress reduction and group cognitive-behavioral stress management on biological markers and psychological symptoms in patients with essential hypertension. This research is a randomized, pretest, post-test, with a control group design, in the form of randomized clinical trial. 60 patients with essential blood pressure who are not controlled their blood pressure by medication will allocate randomly to one of the 3 groups: group of mindfulness-based stress reduction (MBSR), group of cognitive-behavioral stress management and control group. 8 weekly sessions (each session 1/5 hours) group intervention based on MBSR for MBSR group and 8 weekly sessions (each session 1/5 hours) group intervention based on cognitive-behavioral stress management for another group will run. Before and after treatment, research variables such as blood pressure, interleukin 6, oxidative stress, quality of life, anxiety and perceived stress will be assessed and evaluated.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2017103137145N1**
Registration date: **2017-11-14, 1396/08/23**
Registration timing: **prospective**

Last update:

Update count: **0**

Registration date

2017-11-14, 1396/08/23

Registrant information

Name

Farshad Sheybani

Name of organization / entity

Mashhad University of Medical Sciences

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Recruitment status

Recruitment complete

Funding source

This research is related to the PhD dissertation and its funding will be provided by the Iran University of Medical sciences

Expected recruitment start date

2017-11-16, 1396/08/25

Expected recruitment end date

2017-12-11, 1396/09/20

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Comparison the effectiveness of group mindfulness-based stress reduction and group cognitive-behavioral stress management on biological markers and psychological symptoms in patients with essential hypertension

Public title

Effect of stress reduction on blood pressure

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria: Diagnosis of essential hypertension with uncontrolled hypertension despite the treatment according to a cardiologist evaluation; Age between 30 and 60 years; Consent to participate in the research based on the written informed consent form. Exclusion criteria: Secondary blood pressure; Having major depressive disorder; Generalized anxiety disorder; Panic disorder and Post traumatic stress disorder during 1 year ago; Receiving Mindfulness or cognitive-behavioral therapy by a psychologist or psychiatrist in the past.

Age

From **30 years** old to **60 years** old

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **60**

Randomization (investigator's opinion)

Randomized

Randomization description

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

Iran University of Medical Sciences

Street address

Shahid Hemmat Highway, Tehran.

City

Tehran

Postal code

1333813444

Approval date

2017-06-21, 1396/03/31

Ethics committee reference number

IR.IUMS.FMD.REC 1396.9211521215

Health conditions studied

1

Description of health condition studied

Essential hypertension

ICD-10 code

I10

ICD-10 code description

Essential (primary) hypertension

Primary outcomes

1

Description

blood pressure

Timepoint

before and after of intervention

Method of measurement

Home Blood Pressure Monitoring (HBPM)

2

Description

stress oxidative

Timepoint

before and after of intervention

Method of measurement

proxidane and antioxidants balance (PAB)

3

Description

interleukin 6

Timepoint

before and after of intervention

Method of measurement

ELISA method using the Biovendor Germany IL-6 kit

4

Description

anxiety

Timepoint

before and after of intervention

Method of measurement

Beck Anxiety Inventory

5

Description

quality of life

Timepoint

before and after of intervention

Method of measurement

The World Health Organization Quality of Life Instrument
- short form (WHOQOL-BREF)

6

Description

perceived stress

Timepoint

before and after of intervention

Method of measurement

perceived stress scale (PSS) cohen

Secondary outcomes

empty

Intervention groups**1****Description**

The MBSR-based group intervention is based on the 8th session of Jon Kabat Zinn's approach, which lasts 1.5 hours each session.

Category

Lifestyle

2**Description**

Cognitive-behavioral stress management group sessions are based on the Cognitive-Behavioral Stress Management Book (Anthony, Ironson, and Schneiderman, 2010). The duration of each session is 1.5 hours.

Category

Behavior

3**Description**

The control group receives standard blood pressure treatment.

Category

Treatment - Drugs

Recruitment centers**1****Recruitment center****Name of recruitment center**

Rasoul Akram Hospital

Full name of responsible person

Farshad Sheybani

Street address**City**

Tehran

Sponsors / Funding sources**1****Sponsor****Name of organization / entity**

Iran University of Medical Sciences

Full name of responsible person

Dr Behrooz Birashk

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School of Behavioral Sciences and Mental Health,
Mansouri St, Niyayesh St, Sattarkhan St, Tehran, Iran.

City

Tehran

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

Yes

Title of funding source

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

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Position

Ph.D student in clinical psychology

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Other areas of specialty/work

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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