

Clinical Trial Protocol

Iranian Registry of Clinical Trials

28 Jun 2026

Comparison of the Effectiveness of Mindfulness and Body Psychotherapy on Hippocampal Modulation (Volume and Function) and Depression and Executive Function in Patients with Post-Stroke Depression

Protocol summary

Study aim

The main purpose of this study was to find if body psychotherapy and mindfulness interventions that improve the symptoms of depression can change (increase) the hippocampal volume in depressed patients after stroke.

Design

This study is a three-group clinical trial with pre-test and post-test quasi-experimental design. This study has two intervention groups and one control group. The number of patients is 30 women who are randomly divided into three groups.

Settings and conduct

The intervention center is the Anna Mehr Elderly Center. Methods: First, the purposeful sampling will be done based on the patients' scores on the Beck Depression Inventory and the NIHSS test score. It will be taken as a pretest. Patients will be randomly assigned to either intervention or control groups. After the intervention, the questionnaires and computerized tests will be completed again as well as analyzing the recorded scores and comparing the results in the pre-test and post-test stages.

Participants/Inclusion and exclusion criteria

Inclusion criteria: The lesion is located in the left hemisphere; they have a history of stroke only once. Non-inclusion criteria: The patient has a history of a second or more strokes; a history of another chronic illness.

Intervention groups

Intervention group 1: The group that receives the conscious mind. This group includes ten female patients. The intervention is administered to patients during sixteen one-hour sessions, two days a week. Intervention group 2: The second group receives physical therapy intervention. This group includes ten female patients who receive physical therapy intervention for one hour a

day, three days a week. Control group: This group consisted of ten female patients who don't receive any intervention.

Main outcome variables

Depression, executive function, function and volume of hippocamp

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20190812044518N1**

Registration date: **2019-10-08, 1398/07/16**

Registration timing: **retrospective**

Last update: **2019-10-08, 1398/07/16**

Update count: **0**

Registration date

2019-10-08, 1398/07/16

Registrant information

Name

Sahar sadae Nazm Bojnourdi

Name of organization / entity

Ferdowsi of University

Country

Iran (Islamic Republic of)

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+98 58 3222 6225

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2019-02-11, 1397/11/22

Expected recruitment end date

2019-02-21, 1397/12/02

Actual recruitment start date

2019-02-11, 1397/11/22

Actual recruitment end date

2019-03-10, 1397/12/19

Trial completion date

2019-07-15, 1398/04/24

Scientific title

Comparison of the Effectiveness of Mindfulness and Body Psychotherapy on Hippocampal Modulation (Volume and Function) and Depression and Executive Function in Patients with Post-Stroke Depression

Public title

Comparison of the Effectiveness of Mindfulness and Body Psychotherapy on Hippocampus Modulation

Purpose

Supportive

Inclusion/Exclusion criteria**Inclusion criteria:**

Lasting more than three months after a stroke a left hemisphere lesion confirmed by a neurologist and an MRI or CT scan A history of having stroke only for one time ability to read and write depression test score above ten

Exclusion criteria:

The patient has a history of a second stroke Simultaneous use of tobacco and smoking The patient has a history of other chronic illnesses The patient has a history of psychological illness other than depression Patient with aphasia and impaired consciousness The patient has acute economic problems

Age

From **35 years** old to **65 years** old

Gender

Female

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **36**

Actual sample size reached: **30**

Randomization (investigator's opinion)

Randomized

Randomization description

Simple randomization using statistical site

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 Bojnourd University of Medical Sciences

Street address

Bojnourd University of Medical Sciences, Janbaz Blvd, Talghni street

City

Bojnourd

Province

North Khorasan

Postal code

94149-74877

Approval date

2019-08-23, 1398/06/01

Ethics committee reference number

IR.NKUMS.REC.1398.033

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

Post-stroke depression

ICD-10 code**ICD-10 code description****Primary outcomes****1****Description**

Depression score in Beck Questionnaire

Timepoint

Depression test scores are measured before the intervention as pre-test and also in the first intervention group at the end of the eighth week of mindfulness intervention and in the second intervention group at the end of the twelfth week, body psychotherapy intervention.

Method of measurement

Beck Depression Questionnaire

2**Description**

National Institutes of Health(NIH) Stroke Scale Questionnaire Score

Timepoint

National Institutes of Health(NIH) Stroke Scale Questionnaire Score is measured before the intervention as pre-test and also in the first intervention group at the end of the eighth week of mindfulness intervention and in the second intervention group at the end of the twelfth week, body psychotherapy intervention.

Method of measurement

National Institutes of Health(NIH) Stroke Scale Questionnaire

3

Description

Wechsler Digit Span measurement tool Score

Timepoint

Wechsler Digit Span measurement tool Score is measured before the intervention as pre-test and also in the first intervention group at the end of the eighth week of mindfulness intervention and in the second intervention group at the end of the twelfth week, body psychotherapy intervention.

Method of measurement

Wechsler Digit Span measurement tool

4

Description

Course block Computer Psychological Test Score

Timepoint

Course block Computer Psychological Test Score is measured before the intervention as pre-test and also in the first intervention group at the end of the eighth week of mindfulness intervention and in the second intervention group at the end of the twelfth week, body psychotherapy intervention.

Method of measurement

Course block Computer Psychological Test

Secondary outcomes

1

Description

Wisconsin Card Test score

Timepoint

Wisconsin Card Test score is measured before the intervention as pre-test and also in the first intervention group at the end of the eighth week of mindfulness intervention and in the second intervention group at the end of the twelfth week, body psychotherapy intervention.

Method of measurement

Wisconsin Card Test

2

Description

London Tower Test score

Timepoint

London Tower Test score is measured before the intervention as pre-test and also in the first intervention group at the end of the eighth week of mindfulness intervention and in the second intervention group at the end of the twelfth week, body psychotherapy intervention.

Method of measurement

London Tower Test

Intervention groups

1

Description

Intervention group 1: This intervention group receives mindfulness for sixteen one-hour sessions, two days a week. This stress reduction protocol is designed by Kabat Zain (1990). Techniques in this section include relaxation, meditation and meditation along with simple yoga exercises.

Category

Rehabilitation

2

Description

Intervention group 2: This group receives physical therapy intervention for thirty-six one hour sessions, three days a week. This protocol is based on a mix motor and psychological exercises designed by Rohrichet in 2013. Among the techniques in this section are systematic relaxation, balance exercises and rhythmic exercises with breathing and eye movement desensitisation reprocessing.

Category

Rehabilitation

3

Description

Control group: Ten patients who don't receive any intervention in this group.

Category

Other

Recruitment centers

1

Recruitment center

Name of recruitment center

Wellbeing - Mehrana Elderly Center

Full name of responsible person

Dr. Azam Benny Hashim Rad

Street address

Vali Asr 2 alley., Vali Asr street., Farhangian Town

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Ferdowsi University of Mashhad

Full name of responsible person

Dr. Ali Ghanaie Chamanabad

Street address

Ferdowsi University of Mashhad, Azadi Square,

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Province

Razavi Khorasan

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Phone

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Fax**Email**

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

No

Title of funding source

Ferdowsi University of Mashhad

Proportion provided by this source

100

Public or private sector

Private

Domestic or foreign origin

Domestic

Category of foreign source of funding

empty

Country of origin**Type of organization providing the funding**

Persons

Person responsible for general inquiries**Contact****Name of organization / entity**

Mashhad Ferdowsi University

Full name of responsible person

Sahar Sadat Nazm Bojnourdi

Position

Postgraduate student

Latest degree

Bachelor

Other areas of specialty/work

Psychology

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

Yes - There is a plan to make this available

Study Protocol

Yes - There is a plan to make this available

Statistical Analysis Plan

Yes - There is a plan to make this available

Informed Consent Form

Yes - There is a plan to make this available

Clinical Study Report

Yes - There is a plan to make this available

Analytic Code

Yes - There is a plan to make this available

Data Dictionary

Not applicable

Title and more details about the data/document

Data can be shared transparently and without identification variables

When the data will become available and for how long

Start access period one year after publishing the results in a reputable journal

To whom data/document is available

Data will be available only to university researchers and university affiliates

Under which criteria data/document could be used

The data will be available only for the purpose of modeling the process and comparing the results and forecasting the scientific results...

From where data/document is obtainable

Sending e mail to responsible author for access to data
Email sdoctornazmb@gmail.com

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

The data will be made available to the researcher one or two weeks after receiving the email from the applicant.

Comments