

Clinical Trial Protocol

Iranian Registry of Clinical Trials

21 Jul 2026

Investigation the effect of reflexology on fatigue, stress, and cortisol level in women with multiplesclerosis

Protocol summary

Study aim

Determine the effect of reflexology, cortisol stress and cortisol levels in women with multiple sclerosis, and this intervention to improve the quality of life

Design

A clinical trial with a single blind, randomized, control group of 60 women with multiple sclerosis

Settings and conduct

This study is done in multiple sclerosis patients in MS clinic in Yazd, using a one-stop technique with two groups of intervention and control

Participants/Inclusion and exclusion criteria

The population of this study is women with multiple sclerosis of 20 to 40 years old. Entry requirements: history of at least one year of MS, type of relapse, recurrence disorder, disability rating of 1-4 / 5, non compliance: mental illness, acute phase of the disease, corticosteroid use, pregnancy, smoking, alcohol and drugs, participation in yoga classes

Intervention groups

Before and after the intervention and one month after the intervention, the Fatigue Questionnaire and the stress test were completed and interviewed by a neurologist and psychologist. Cortisol was taken in the intervention group for four weeks, three times a week for foot reflexology. First, she is trained by the researcher and in the presence of her, and she is trained to do this at home three times a week and is conducted by the researcher for telephone massage. The patient is asked to lie down the bed in bed. The foot reflexology was then resting for 30 minutes (for 15 minutes each) and directing the Solar Plexus at 6 minutes, 3 minutes, and again for 6 minutes, at a pressure of rotational pressure of 4-3 Kg, weekly 3 sessions for 4 weeks. In the control group, creating the same conditions with the intervention group, simple foot massage of the foot without compression of the standard reflexology points of the foot is performed with the same duration of the intervention group

Main outcome variables

Fatigue score, estress score, cortisol level

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20180304038936N1**

Registration date: **2018-07-10, 1397/04/19**

Registration timing: **registered_while_recruiting**

Last update: **2018-07-10, 1397/04/19**

Update count: **0**

Registration date

2018-07-10, 1397/04/19

Registrant information

Name

Ahmadreza Sayadi

Name of organization / entity

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2018-06-24, 1397/04/03

Expected recruitment end date

2018-09-25, 1397/07/03

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Investigation the effect of reflexology on fatigue, stress, and cortisol level in women with multiplesclerosis

Public title

Investigation the effect of reflexology on fatigue, stress and cortisol level in women with multiple sclerosis

Purpose

Supportive

Inclusion/Exclusion criteria**Inclusion criteria:**

The absence of the patient in the acute phase of the disease Age 20 to 40 years old Type of relapse-suppression disease Having astrong companion Disability score rating 1-4/5 Iranian nationality

Exclusion criteria:

Simultaneous physical illness Mental illness Acute stage of the disease corticosteroid use two months befor study Pregnancy Smoking,alchol and narcotics Participating in yoga classes, meditation and relaxation Hearing impairment and tongue Lesion on the foot of the foot

Age

From **20 years** old to **40 years** old

Gender

Female

Phase

N/A

Groups that have been masked

- Participant

Sample size

Target sample size: **60**

Randomization (investigator's opinion)

Randomized

Randomization description

In order to randomize this study, the first two groups are selected by lottery method.Selecting the rest of the group is done by minimizing the procedure.The tool is a randomized packet

Blinding (investigator's opinion)

Single blinded

Blinding description

Participants do not know which group is being investigated as intervention group.

Placebo

Used

Assignment

Parallel

Other design features**Secondary Ids**

empty

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

Ethics committee of Rafsangan University of Medical Sciences

Street address

Central Office of Medical Sciences, imam ali Blvd, Rafsangan

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Province

Kerman

Postal code

771793777

Approval date

2018-04-17, 1397/01/28

Ethics committee reference number

IR.RUMS.REC.1397.026

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

Fatigue

ICD-10 code**ICD-10 code description****2****Description of health condition studied**

Stress

ICD-10 code**ICD-10 code description****3****Description of health condition studied**

Cortisol

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

Fatigue

Timepoint

Befor the intervention, one month after the intervention, one month after the end of intervention

Method of measurement

Fatigue score is obtained using the fatigue criterion.this tool consists of 9 questions with a grading scale of 1-7.

2**Description**

Stress

Timepoint

Befor the intervention, one month after the intervention, one month after the end of intervention

Method of measurement

Stress score is obtained by stress syndrome

questionnaire. The questionnaire consists of 50 items that are based on a 6-point Likert scale, and measures four subscales of physical, emotional, cognitive, and behavioral.

3

Description

cortisol

Timepoint

Before the intervention, one month after the intervention, one month after the end of intervention

Method of measurement

At 8 o'clock in the morning, take the patient's blood sample and it will be sent to the laboratory within one to one hour.

Secondary outcomes

1

Description

Fatigue rating based on fatigue severity questionnaire

Timepoint

Before the intervention, one month after the intervention, one month after the end of the intervention

Method of measurement

The questionnaire was used to measure the fatigue intensity of physical, psychological, emotional, behavioral, and social dimensions. This tool consists of nine questions ranging from one to seven. Exhaustion is considered to be a mild fatigue, moderate fatigue, severe fatigue.

2

Description

Stress score based on stress syndrome questionnaire

Timepoint

Before the intervention, one month after the intervention, one month after the end of the intervention

Method of measurement

Stress syndrome questionnaire consists of 4 subscales of physical, emotional, cognitive and therapeutic exercises and 50 items. Scoring this scale is based on a 6-point Likert scale

3

Description

cortisol level

Timepoint

Before the intervention, one month after the intervention, one month after the end of the intervention

Method of measurement

Patients' blood samples are taken from the patient at the time indicated, and then sent to the laboratory at -20 ° C after centrifugation

Intervention groups

1

Description

Intervention group: The patient is asked to lie down the bed in bed. Then, we perform foot reflexology for 15 minutes per leg.

Category

N/A

Recruitment centers

1

Recruitment center

Name of recruitment center

MS Society located at Shahid Sadoughi Hospital, Yazd

Full name of responsible person

Ahmad reza sayadi

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

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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

Rafsanjan 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

Rafsanjan University of Medical Sciences

Full name of responsible person

Ahmad reza sayadi

Position

ph.d

Latest degree

Ph.D.

Other areas of specialty/work

Nursery

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Sharing plan

Deidentified Individual Participant Data Set (IPD)

Undecided - It is not yet known if there will be 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

Undecided - It is not yet known if there will be 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