

Clinical Trial Protocol

Iranian Registry of Clinical Trials

09 Jul 2026

Investigating the effectiveness of the acupressure combination of Sanyinjiao (SP6) and Shenmen (HT7) points on the sleep quality of the elderly: A family-oriented three parallel group clinical trial

Protocol summary

Study aim

Determining the effectiveness of the acupressure combination of Sanyinjiao and Shenmen points on the sleep quality of the elderly

Design

Clinical trial of three groups without control group, with parallel groups, randomized, without blinding

Settings and conduct

The location of the study will be the clinics of Shahroud city. In a face-to-face meeting, the researcher teaches the elderly caregiver how to press the mentioned points, and they must perform this action on the elderly once a day at night time at 10-11 pm. Before and one week after the intervention, Pittsburgh questionnaire will be completed for all three groups.

Participants/Inclusion and exclusion criteria

Entering criteria for elderly: age > 60 years, informed consent, MMSE>22 in literate people and AMTS>7 in illiterate people, unfavorable sleep quality based on Pittsburgh questionnaire. Exclusion criteria for elderly: receiving other sleep-related treatments within 2 weeks before the study, history of mental illness, dependence or addiction to any narcotic, painkiller, antidepressant. Inclusion criteria for the elderly caregiver: having the most contact with the elderly, the health literacy score more than 66.1 from the HELIA questionnaire. Exclusion criteria for the elderly caregiver: Having any physical disease related to the fingers

Intervention groups

In group A (Shenmen), B (Sanyinjiao) and C (combination of Shenmen and Sanyinjiao), in a face-to-face meeting, the researcher will teach the person accompanying the elderly to press the mentioned points, and they should do it once a day for a week at 10-11 pm, for elderly. This study has no control group.

Main outcome variables

Sleep quality

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20230222057494N1**

Registration date: **2023-04-05, 1402/01/16**

Registration timing: **prospective**

Last update: **2023-04-05, 1402/01/16**

Update count: **0**

Registration date

2023-04-05, 1402/01/16

Registrant information

Name

Reza Mirafzali

Name of organization / entity

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2023-04-13, 1402/01/24

Expected recruitment end date

2023-11-11, 1402/08/20

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Investigating the effectiveness of the acupressure combination of Sanyinjiao (SP6) and Shenmen (HT7) points on the sleep quality of the elderly: A family-oriented three parallel group clinical trial

Public title

Acupressure combination of Sanyinjiao (SP6) and Shenmen (HT7) points and sleep quality of the elderly

Purpose

Treatment

Inclusion/Exclusion criteria**Inclusion criteria:**

age over 60 years old Having full awareness and listening and speaking ability suitable for learning research intervention MMSE score more than 22 in literate people and AMTS score more than 7 in illiterate people Unfavorable sleep quality based on answers to the Pittsburgh sleep quality questionnaire (a score of 5-21 is considered as unfavorable sleep quality) Entry criteria for the person accompanying the elderly: 1- To have the most contact with the elderly. To have the ability to correctly implement the intervention on the elderly The health literacy score is more than 66.1 from the HELIA questionnaire

Exclusion criteria:

Secondary insomnia caused by physical illness or mental disorder Those who received other sleep-related treatments within 2 weeks before the study. History of mental illness (based on the elderly's self-report) Dependence or addiction to any drug, pain reliever, anti-depressant, sleeping pill and alcohol Having sleep-related diseases such as obstructive sleep apnea or restless legs syndrome Regarding the person accompanying the elderly: Having any physical disease related to the fingers

Age

From **60 years** old

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **126**

Randomization (investigator's opinion)

Randomized

Randomization description

Patients will be divided into three groups through randomization and based on the production of random numbers strings through software. (www.randomizer.org) In order for the number of assignments to each group to be equal, the block method of volume 6 is used to create a random allocation sequence because participants are gradually entered into the study. The allocation concealment will be done centrally. In this way, the assignment sequence will be done by someone outside the research team. Upon each person entering the study, the researcher will contact the person and assign the person to the group.

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 Shahroud University of Medical Sciences

Street address

Shahroud University of Medical Sciences, Hafte-Tir Sq., Tehran Street

City

Shahroud

Province

Semnan

Postal code

3614773955

Approval date

2023-02-05, 1401/11/16

Ethics committee reference number

IR.SHMU.REC.1401.219

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

Sleep quality

ICD-10 code

G47.9

ICD-10 code description

Sleep disorder, unspecified

Primary outcomes**1****Description**

Sleep Quality Based on Pittsburgh Sleep Quality Test Score

Timepoint

Before and one week after the intervention

Method of measurement

Pittsburgh Sleep Quality Questionnaire

Secondary outcomes

empty

Intervention groups

1

Description

Intervention group 1: In this group (group A), first in a face-to-face session, pressing the Shenmen point (HT7) (on the ulnar side, the distal volar bands of the wrist and the radial border of the flexor carpi ulnaris tendon in the anterior region of the wrist) with A pressure of 3 to 4 kg using a circular motion with a speed of two revolutions per second in a firm and uniform way for one minute pressure 30 seconds rest up to 3 times with the thumb by the researcher is taught to the person with the intervention group and they must do it once every day at night at 10-11 for a week. Perform this procedure on the elderly themselves and repeat it.

Category

Treatment - Other

2

Description

Intervention group 2: In this group (group B), first, in a face-to-face session, pressing the Sanyinjiao point (SP6) (directly to the tip of the medial malleolus, the posterior margin of the medial leg of the leg) with a pressure of 3 to 4 kg using A circular movement with a speed of two revolutions per second in a firm and uniform way for one minute, pressure for 30 seconds, rest up to 3 times with the thumb is taught by the researcher to the person accompanying the people of the intervention group, and they should do this procedure on the elderly themselves for a week, once every day at night time at 10-11 and repeat.

Category

Treatment - Other

3

Description

Intervention group 3: In this group (group C), first in a face-to-face meeting, pressing the Shenmen point (HT7) (on the ulnar side, the distal volar bands of the wrist and the radial border of the flexor carpi ulnaris tendon in the anterior region of the wrist) and the Sanyinjiao point (SP6) (straight up to the tip of the medial malleolus, the posterior border of the inner leg of the leg) with a pressure of 3 to 4 kg using a circular motion at a speed of two rotations per second in a firm and uniform manner for one minute of pressure and 30 seconds of rest. Up to 3 times, the person accompanying the intervention group will be trained by the researcher with the thumb up to 3 times, and they They should perform and repeat this procedure on the elderly once every day at night at 10-11 for a week and repeat.

Category

Treatment - Other

Recruitment centers

1

Recruitment center

Name of recruitment center

Clinics in Shahroud city

Full name of responsible person

Dr. Hossein Bagheri

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Shahroud University of Medical Sciences

Full name of responsible person

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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

Shahroud 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

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Person responsible for general inquiries

Contact

Name of organization / entity

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Full name of responsible person

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Position

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Latest degree

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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