

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

10 Jun 2026

### Effectiveness of Cognitive-Based Occupational Therapy Interventions on The Level of Cognitive Performance of Elderly Women with Mild Cognitive Impairment (MCI) In Shiraz City: A Randomized Controlled Trial

#### Protocol summary

##### Study aim

Determining the effect of cognitive-based occupational therapy interventions on cognitive function of elderly women with mild cognitive impairment (MCI) in Shiraz city

##### Design

A randomized clinical trial using the Ephron equal block randomization method, in a single-blind manner, the sample size was obtained using NCCS-PASS version 15 software, 62 people, who were equally allocated into 2 groups (intervention and control).

##### Settings and conduct

Intervention classes and questionnaire completion are conducted in the Farzanegan Foundation of Shiraz in the presence of the facilitator and the clinic manager. Blinding occurs in the participants in such a way that they are unaware of the study group allocation and all participants involved in the study assume that all participants have received the intervention.

##### Participants/Inclusion and exclusion criteria

Inclusion criteria: Elderly women with mild cognitive impairment evaluated with the MoCA test with a score of less than 26, Elderly women with memory complaints, Willingness to participate in the study Exclusion criteria: Having more than one absent session or more in treatment programs, Elderly women who attended the pre-test but did not attend the post-test, Incomplete completion of questionnaires, Transfer of residence or long-term illness or refusal of treatment and death.

##### Intervention groups

The elderly will be divided into two groups: intervention and control, and the intervention group will receive a class to increase cognitive skills. The control group will receive yoga classes from Soroush Center.

##### Main outcome variables

Mild cognitive impairment, executive functions, cognitive flexibility, memory, cognitive-based occupational therapy

interventions

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20180514039648N9**

Registration date: **2024-12-25, 1403/10/05**

Registration timing: **prospective**

Last update: **2024-12-25, 1403/10/05**

Update count: **0**

##### Registration date

2024-12-25, 1403/10/05

##### Registrant information

##### Name

Abdolrahim Asadollahi

##### Name of organization / entity

##### Country

Iran (Islamic Republic of)

##### Phone

+98 71 3725 1001

##### Email address

a\_asadollahi@sums.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2025-01-08, 1403/10/19

##### Expected recruitment end date

2025-04-08, 1404/01/19

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

## Trial completion date

empty

## Scientific title

Effectiveness of Cognitive-Based Occupational Therapy Interventions on The Level of Cognitive Performance of Elderly Women with Mild Cognitive Impairment (MCI) In Shiraz City: A Randomized Controlled Trial

## Public title

Effectiveness of Cognitive-Based Occupational Therapy Interventions on The Level of Cognitive Performance of Elderly Women with Mild Cognitive Impairment (MCI)

## Purpose

Prevention

## Inclusion/Exclusion criteria

### Inclusion criteria:

Elderly women with mild cognitive impairment evaluated with the MoCA test with a score less than 26 Elderly women with memory complaints Willingness to participate in the study

### Exclusion criteria:

Having more than one absent session or more in treatment programs Elderly women who attended the pre-test but did not attend the post-test. Incomplete completion of questionnaires Transfer of residence or long-term illness or refusal of treatment and death.

## Age

From **60 years** old

## Gender

Female

## Phase

3

## Groups that have been masked

- Participant

## Sample size

Target sample size: **62**

## Randomization (investigator's opinion)

Randomized

## Randomization description

In the intervention phase in random trials, for sampling the study samples into different groups of control and control groups, randomly divided in this type of trials, a group of people who are members of Farzangan Shiraz Foundation and meet the entry criteria will be selected (Random Allocation) In each care center for the elderly, there is a list of the names of the members in the center with the national code, and each number will be defined as a variable ID number in the software. We will select about 200 members of Farzangan Shiraz Foundation and measure them using the Montreal Cognitive Status Questionnaire (MOCA) and 62 people who meet all the conditions for entering the study will be selected and using the technique Random sampling in such a way that these 31 people will be classified into 2 equal blocks in each block of 31 people between the groups and the final sample will be set. Random allocation of elderly people referring to the day care center for the elderly with the help of statistical software in intervention and control groups with Efron's randomization algorithm with blocks equal to Efron's  $P = 0.75$  and 500 rotations (chance in

the control and intervention groups is done using blinding In this Procedures DOE► Randomization Lists version 15.0.1 with the command CSS-PASS, the study was a blind study so that the elderly are unaware of their group title.

## Blinding (investigator's opinion)

Single blinded

## Blinding description

Blinding occurs in participants in such a way that they are unaware of the study group allocation and all participants involved in the study assume that all participants have received the intervention.

## Placebo

Not used

## Assignment

Parallel

## Other design features

## Secondary Ids

empty

## Ethics committees

### 1

#### Ethics committee

##### Name of ethics committee

Ethics Committee of Shiraz University of Medical Sciences

##### Street address

School of Health, Razi Ave.

##### City

Shiraz

##### Province

Fars

##### Postal code

7153675541

#### Approval date

2024-12-11, 1403/09/21

#### Ethics committee reference number

IR.SUMS.SCHEANUT.REC.1403.084

## Health conditions studied

### 1

#### Description of health condition studied

Mild cognitive impairment

#### ICD-10 code

G31. 84

#### ICD-10 code description

Mild cognitive impairment of uncertain or unknown etiology

## Primary outcomes

### 1

#### Description

Executive Functions

#### Timepoint

Before the intervention, immediately after the end of the intervention, and two months after the intervention

**Method of measurement**

Tower of London test

**2**

**Description**

Cognitive flexibility

**Timepoint**

Before the intervention, immediately after the end of the intervention, and two months after the intervention

**Method of measurement**

Wisconsin Card Sorting Inspired Task (WCST)

**3**

**Description**

memory

**Timepoint**

Before the intervention, immediately after the end of the intervention, and two months after the intervention

**Method of measurement**

Wechsler Memory Scale

**Secondary outcomes**

empty

**Intervention groups**

**1**

**Description**

Intervention group: Intervention group: 12 one-hour intervention sessions will be held for the intervention group (31 people) with the aim of increasing the cognitive skills of elderly women. In the intervention group, the Wechsler Memory Inventory and the Wisconsin Card Test along with the Tower of London Test will be administered before the start of the intervention, immediately after the intervention, and 2 months after the end of the intervention.

**Category**

Prevention

**2**

**Description**

Control group: Control group: 12 one-hour yoga and exercise class intervention sessions will be conducted for the control group at the Soroush older adults Center. In the control group, the Wechsler Memory Inventory and the Wisconsin Card Test along with the Tower of London Test will be administered before the start of the intervention, immediately after the intervention, and 2 months after the end of the intervention. After completing the intervention package provided to the experimental group, it will also be provided to the control group.

**Category**

Prevention

**Recruitment centers**

**1**

**Recruitment center**

**Name of recruitment center**

FDCF of Shiraz

**Full name of responsible person**

Dr.Abdolrahim Asadollahi

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No . 30, Dinakan5 , Mirzaye Shirazi Blvd, Shiraz

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**Sponsors / Funding sources**

**1**

**Sponsor**

**Name of organization / entity**

Shiraz 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**

Shiraz 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**  
Shiraz University of Medical Sciences

**Full name of responsible person**  
Mahdis Ferasat

**Position**  
Trial Project Associate and Student

**Latest degree**  
Master

**Other areas of specialty/work**  
Geriatrics

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## Person responsible for scientific inquiries

### Contact

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## Person responsible for updating data

### Contact

**Name of organization / entity**

Shiraz University of Medical Sciences

### Full name of responsible person

Abdolrahim Asadollahi

### Position

Associate professor

### Latest degree

Ph.D.

### Other areas of specialty/work

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## Sharing plan

### Deidentified Individual Participant Data Set (IPD)

Yes - There is a plan to make this available

### 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

Yes - There is a plan to make this available

### Analytic Code

No - There is not a plan to make this available

### Data Dictionary

Not applicable

### Title and more details about the data/document

The datasets generated and/or analysed during the current study are available from the authors upon reasonable request and with the permission of SUMS.

### When the data will become available and for how long

Immediately at the time of printing the results

### To whom data/document is available

Data will be made available to academic researchers upon request.

### Under which criteria data/document could be used

Allowed with source reference.

### From where data/document is obtainable

Scientific supervisor of the study: Dr. Abdolrahim Asadollahi by sending him an official email.

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

Upon receiving an email from the applicant and reviewing his/her academic qualifications, data will be shared immediately to the same email.

### Comments