

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

08 Jul 2026

### study the effect of one shot cognitive behavioral therapy on insomnia and heart rate variability of healthcare workers at the time of covid-19

#### Protocol summary

insomnia, heart rate variability

#### Study aim

Study the effect of one shot cognitive behavioral therapy on insomnia and heart rate variability of health care workers at the time of Covid-19 pandemic

#### Design

Clinical trial, with parallel groups, for 70 patients, randomized using a random string, double-blind, the outcome evaluator and data analyzer will be blinded,

#### Settings and conduct

This study is being performed on health care workers at Al-Zahra and Khorshid Hospitals in Isfahan. 70 healthcare workers will randomly divide into two groups and the intervention group underwent a one-shot cognitive behavioral therapy intervention. Blinding is done by assigning a unique code to patients and the outcome assessor and the data analyzer are unaware of the groups.

#### Participants/Inclusion and exclusion criteria

Inclusion Criteria: Healthcare workers who work in ALzahra hospital, insomnia score of 8 and higher based on Insomnia Severity Index, age 18 years old and above; Exclusion criteria: History of cardiovascular disease, history of seizure, manic depressive disorder, narcolepsy, and parasomnia, using beta-adrenergic blockade or antiarrhythmic medications

#### Intervention groups

Intervention group: The sleep diary sheets are given to participants to be completed by them within the next 2 weeks. Then participants are divided into groups of 2-3 people and each group will come for a group meeting (for one-shot cognitive behavioral therapy) at a specific time. An ECG will be recorded for 5 minutes before the start of the session to assess changes in basal heart rate. After a month, the consequences are reviewed. Control group: includes hospital health staff for whom no intervention is performed and only at similar times in the intervention group, heart rate changes and insomnia severity are evaluated for them.

#### Main outcome variables

#### General information

##### Reason for update

This study is carried on healthcare workers. During the recruitment, they were involved with the 4th wave of the Covid-19 pandemic. Hence, we had to change the protocol.

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20171219037964N4**

Registration date: **2021-03-20, 1399/12/30**

Registration timing: **prospective**

Last update: **2021-12-18, 1400/09/27**

Update count: **1**

##### Registration date

2021-03-20, 1399/12/30

##### Registrant information

###### Name

Babak Amra

###### Name of organization / entity

###### Country

Iran (Islamic Republic of)

###### Phone

+98 31 3668 0048

###### Email address

amra@med.mui.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2021-04-09, 1400/01/20

##### Expected recruitment end date

2021-07-22, 1400/04/31

##### Actual recruitment start date

2021-04-21, 1400/02/01  
**Actual recruitment end date**  
2021-07-22, 1400/04/31  
**Trial completion date**  
2021-09-22, 1400/06/31

**Scientific title**  
study the effect of one shot cognitive behavioral therapy on insomnia and heart rate variability of healthcare workers at the time of covid-19

**Public title**  
study the effect of cognitive therapy on insomnia and heart rate variability of health care workers

**Purpose**  
Treatment

**Inclusion/Exclusion criteria**

**Inclusion criteria:**  
Healthcare workers who work in ALzahra and Khorshid hospital Insomnia score of 8 and higher based on insomnia severity index age over 18 years old

**Exclusion criteria:**  
History of cardiovascular disease History of seizure, manic depressive disorder, narcolepsy and parasomnia Using beta adrenergic blockade or anti arrhythmic medications

**Age**  
From **18 years** old

**Gender**  
Both

**Phase**  
N/A

**Groups that have been masked**

- Care provider
- Investigator
- Outcome assessor
- Data analyser

**Sample size**  
Target sample size: **70**  
Actual sample size reached: **57**

**Randomization (investigator's opinion)**  
Randomized

**Randomization description**  
A random sequence for 70 numbers is made using randomization.com and is available to the research assistant. Each person is given a code (from one to 70). The research assistant places the person in the relevant groups by matching the person's code with a random sequence.

**Blinding (investigator's opinion)**  
Double blinded

**Blinding description**  
Each person is given a unique code and these codes are given to the research assistant. The person in charge of data analysis and the person in charge of follow-up are unaware of their allocation to study groups and study codes.

**Placebo**  
Not used

**Assignment**  
Parallel

**Other design features**

**Secondary Ids**

empty

**Ethics committees**

**1**

**Ethics committee**

**Name of ethics committee**

Ethics Committee of Isfahan University of Medical Sciences

**Street address**

Hezarjerib St.

**City**

Isfahan

**Province**

Isfahan

**Postal code**

8174673461

**Approval date**

2021-03-15, 1399/12/25

**Ethics committee reference number**

IR.MUI.MED.REC.1399.1153

**Health conditions studied**

**1**

**Description of health condition studied**

Insomnia

**ICD-10 code**

F51.01

**ICD-10 code description**

Primary insomnia

**Primary outcomes**

**1**

**Description**

insomnia

**Timepoint**

at baseline, after one month

**Method of measurement**

Insomnia Severity Index

**2**

**Description**

Heart rate variability

**Timepoint**

at baseline, after one month

**Method of measurement**

heart monitoring

**Secondary outcomes**

empty

## Intervention groups

### 1

#### Description

Intervention group: This group first completes sleep diary for two weeks and then receives a group session (in groups of 2-3) of a one-session cognitive behavioral intervention based on their sleep status and sleep habits.

#### Category

Treatment - Other

### 2

#### Description

Control group: Routine treatments

#### Category

N/A

## Recruitment centers

### 1

#### Recruitment center

##### Name of recruitment center

Alzahra hospital

##### Full name of responsible person

Babak Amra

##### Street address

Soffe St.

##### City

Isfahan

##### Province

Isfahan

##### Postal code

8174675731

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+98 31 3620 2020

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amra@med.mui.ac.ir

### 2

#### Recruitment center

##### Name of recruitment center

Khorshid Hospital

##### Full name of responsible person

Babak Amra

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Ostandari St.

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8145831451

##### Phone

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## Sponsors / Funding sources

### 1

#### Sponsor

##### Name of organization / entity

Esfahan University of Medical Sciences

##### Full name of responsible person

Shaghayegh Haghjoo

##### Street address

Chancellory of Research, Isfahan University of Medical Sciences, Hezarjerib St.

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sh\_haghjoo@med.mui.ac.ir

#### Grant name

#### Grant code / Reference number

#### Is the source of funding the same sponsor organization/entity?

Yes

#### Title of funding source

Esfahan 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

Esfahan University of Medical Sciences

##### Full name of responsible person

Babak Amra

##### Position

Professor

##### Latest degree

Subspecialist

##### Other areas of specialty/work

Internal Medicine

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##### Province

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## Person responsible for scientific inquiries

### Contact

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**Full name of responsible person**  
Babak Amra  
**Position**  
Professor  
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**Other areas of specialty/work**  
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## Person responsible for updating data

### Contact

**Name of organization / entity**  
Esfahan University of Medical Sciences  
**Full name of responsible person**  
Babak Amra  
**Position**  
Professor  
**Latest degree**  
Subspecialist  
**Other areas of specialty/work**  
Internal Medicine

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**Fax**  
**Email**  
Amra@med.mui.ac.ir

## 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

Yes - There is a plan to make this available

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

Participants data will be Shared after being unidentified

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

Six months after the publication of the results

### To whom data/document is available

All researchers are allowed to have access to data

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

In order to use in Meta-analysis studies

### From where data/document is obtainable

A request should be sent to the Chancellery of Research of Isfahan University of Medical Sciences

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

The request will be reviewed in the Chancellery of Research and the Ethics Committee of Isfahan University of Medical Sciences; Data will be delivered upon approval.

### Comments