

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

24 Jun 2026

### Effectiveness of unified protocol of transdiagnostic treatment on psychological symptoms, fatigue, emotion regulation, health anxiety, sleep quality, and sexual satisfaction in patients with multiple sclerosis

#### Protocol summary

##### Study aim

Present study aims to investigate the effectiveness of unified protocol of transdiagnostic treatment (UP) on psychological symptoms, fatigue, emotion regulation, health anxiety, sleep quality, and sexual satisfaction in patients with multiple sclerosis (MS).

##### Design

This randomized and single-blind clinical trial with parallel and control groups will be conducted on 48 patients who will be randomly selected using the simple random.

##### Settings and conduct

Patients referring to Moheb Mehr Hospital Neurology Clinic, Tehran, Iran are chosen as the participants of the study. In this single-blind study, sealed opaque envelopes will be used to conceal the sequencing. Intervention group will receive 12 sessions of online psychotherapy based on unified protocol of transdiagnostic treatment and control group will not receive the treatment intervention. The responsible for data collection is blind to group allocation and the type of intervention.

##### Participants/Inclusion and exclusion criteria

Statistical society of the study includes all of the MS patients in Tehran, and selected sample will be 48 patients referred to neurology clinic of Moheb Mehr Hospital. Main inclusion criteria include confirmation of diagnosis of MS, being at least 20 y/o, being sexually active, fixed dosage of medicine since 3 month before the beginning of treatment, and not having severe psychiatric and physical illnesses.

##### Intervention groups

Intervention group will receive 12 sessions of online psychotherapy based on unified protocol of transdiagnostic treatment and control group will not receive the treatment intervention.

##### Main outcome variables

Improving psychological symptoms (depression, anxiety, stress), fatigue, emotion regulation, health anxiety, sleep quality, and sexual satisfaction in MS patients.

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20221230056986N1**

Registration date: **2023-06-17, 1402/03/27**

Registration timing: **prospective**

Last update: **2023-06-17, 1402/03/27**

Update count: **0**

##### Registration date

2023-06-17, 1402/03/27

##### Registrant information

##### Name

Amir Mahdi Katani

##### Name of organization / entity

##### Country

Iran (Islamic Republic of)

##### Phone

+98 21 2303 1548

##### Email address

katani@sbmu.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2023-08-23, 1402/06/01

##### Expected recruitment end date

2024-08-22, 1403/06/01

##### Actual recruitment start date

empty

**Actual recruitment end date**

empty

**Trial completion date**

empty

**Scientific title**

Effectiveness of unified protocol of transdiagnostic treatment on psychological symptoms, fatigue, emotion regulation, health anxiety, sleep quality, and sexual satisfaction in patients with multiple sclerosis

**Public title**

Effectiveness of transdiagnostic treatment on psychological parameters in multiple sclerosis

**Purpose**

Treatment

**Inclusion/Exclusion criteria****Inclusion criteria:**

Confirmation of relapsing-remitting multiple sclerosis diagnosis by neurologist Age least 20 years Being sexually active Mastery in Persian language and reading and writing literacy Fixed dosage of medicine from 3 months before beginning the treatment Being physically able to participate in weekly sessions Not being diagnosed with severe psychiatric disorders Not being diagnosed with another serious or chronic physical illness Not having the history of receiving psychological treatments

**Exclusion criteria:****Age**

From 20 years old

**Gender**

Both

**Phase**

2-3

**Groups that have been masked**

- Participant

**Sample size**

Target sample size: 48

**Randomization (investigator's opinion)**

Randomized

**Randomization description**

The process of random allocation of study participants to two groups; intervention group and control group included both creating a random sequence and concealing the random allocation. Firstly, using the random sequence rule, 48 papers for participants of the intervention and control groups (which are marked with A and B and are not defined to avoid bias) were thrown into a lottery container and then the papers were removed from the lottery container -without replacement- and the created sequence is recorded. Then, using the snooze method, 48 envelopes were prepared and each of the random sequences created is recorded on a card and placed in the envelopes in order, and the envelopes were also numbered. Then, the envelopes were given to the participants based on the order of entering the study, so the grouping of each was determined.

**Blinding (investigator's opinion)**

Single blinded

**Blinding description**

Patients referring to Moheb Mehr Hospital Neurology Clinic, Tehran, Iran are chosen as the participants of the study. In this single-blind study, sealed opaque envelopes will be used to conceal the sequencing. Intervention group will receive 12 sessions of online psychotherapy based on unified protocol of transdiagnostic treatment and control group will not receive the treatment intervention. The responsible for data collection is blind to group allocation and the type of intervention.

**Placebo**

Not used

**Assignment**

Parallel

**Other design features****Secondary Ids**

empty

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

Ethics committee of Shahid Beheshti University of Medical Sciences

**Street address**

Shahid Beheshti University of Medical Sciences, Yaman St. Velenjak

**City**

Tehran

**Province**

Tehran

**Postal code**

1985717443

**Approval date**

2019-09-24, 1398/07/02

**Ethics committee reference number**

IR.SBMU.MSP.REC.1398.669

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

Multiple Sclerosis

**ICD-10 code**

G35

**ICD-10 code description**

Multiple sclerosis

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

Psychological symptoms

**Timepoint**

Before intervention, after intervention, three month after

intervention

#### **Method of measurement**

Depression, Anxiety, Stress Scale (DASS-21)

### **2**

#### **Description**

Fatigue

#### **Timepoint**

Before intervention, after intervention, three month after intervention

#### **Method of measurement**

Fatigue Severity Scale (FSS)

### **3**

#### **Description**

Quality of sleep

#### **Timepoint**

Before intervention, after intervention, three month after intervention

#### **Method of measurement**

Pittsburgh Sleep Quality Index (PSQI)

### **4**

#### **Description**

Sexual Satisfaction

#### **Timepoint**

Before intervention, after intervention, three month after intervention

#### **Method of measurement**

Larson Sexual Satisfaction Questionnaire (LSSQ)

### **5**

#### **Description**

Emotion regulation

#### **Timepoint**

Before intervention, after intervention, three month after intervention

#### **Method of measurement**

Cognitive Emotion Regulation Questionnaire (SERQ)

### **6**

#### **Description**

Health anxiety

#### **Timepoint**

Before intervention, after intervention, three month after intervention

#### **Method of measurement**

Health Anxiety Questionnaire (HAQ)

## **Secondary outcomes**

empty

## **Intervention groups**

### **1**

#### **Description**

Intervention group: 12 sessions of online psychotherapy based on unified protocol of transdiagnostic treatment

#### **Category**

Treatment - Other

### **2**

#### **Description**

Control group: Not receiving treatment intervention

#### **Category**

Treatment - Other

## **Recruitment centers**

### **1**

#### **Recruitment center**

##### **Name of recruitment center**

Moheb Mehr Hospital Neurology Clinic

##### **Full name of responsible person**

Reza Hajmanouchehri

##### **Street address**

Khalilzadeh alley, after Vanak square, Valiasr St.

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تهران

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##### **Postal code**

1319919697

##### **Phone**

+98 21 85555

##### **Email**

info@moheb.com

##### **Web page address**

<https://www.moheb.com/mohebmehr/home/>

## **Sponsors / Funding sources**

### **1**

#### **Sponsor**

##### **Name of organization / entity**

Shahid Beheshti University of Medical Sciences

##### **Full name of responsible person**

Afshin Zarghi

##### **Street address**

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##### **Phone**

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##### **Email**

Mpajouhesh@sbmu.ac.ir

##### **Web page address**

<https://research.sbmu.ac.ir/index.jsp?pageid=2554&p=1>

#### **Grant name**

**Grant code / Reference number**  
**Is the source of funding the same sponsor organization/entity?**  
Yes  
**Title of funding source**  
Shahid Beheshti 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**  
Shahid Beheshti University of Medical Sciences  
**Full name of responsible person**  
Amir Mahdi Katani  
**Position**  
Phd Student  
**Latest degree**  
Ph.D.  
**Other areas of specialty/work**  
Psychology  
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amir.katani.1993@gmail.com

## Person responsible for scientific inquiries

### Contact

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## Person responsible for updating data

### Contact

**Name of organization / entity**  
Shahid Beheshti University of Medical Sciences  
**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)

Yes - There is a plan to make this available

### Study Protocol

Yes - There is a plan to make this available

### Statistical Analysis Plan

Not applicable

### Informed Consent Form

Yes - There is a plan to make this available

### Clinical Study Report

Not applicable

### Analytic Code

Not applicable

### Data Dictionary

Not applicable

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

SPSS file of participants data, treatment protocol of study, and informed consent form will be available if necessary.

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

After execution of study

### To whom data/document is available

Editor in chief of scientific journals

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

At request of editor in chief of scientific journals

**From where data/document is obtainable**

By email: amir.katani.1993@gmail.com

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

Data will be sent by email after request.

**Comments**