

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

10 Jun 2026

### Exploring the impact of the Adaptive Disclosure technique on reducing the severity of post-traumatic stress disorder symptoms in veterans: a randomized controlled trial

#### Protocol summary

##### Study aim

The impact of the adaptive disclosure technique on the severity of post-traumatic stress disorder symptoms in veterans with psychological injuries.

##### Design

Block randomization will be conducted with a total sample size of 58 participants, consisting of two groups: intervention and control, with 29 participants in each group.

##### Settings and conduct

This study will be conducted over a period of 6 weeks at Salman Hospital and Shahid Rajaei Hospital in Yasuj, Iran.

##### Participants/Inclusion and exclusion criteria

Inclusion criteria include veterans with psychological injuries diagnosed with post-traumatic stress disorder (co-occurring disorders such as depression, anxiety, or substance abuse are permitted if treated or if there are no ongoing dependency issues). Exclusion criteria include individuals with severe Axis I psychiatric disorders, such as psychotic disorders or Type I bipolar disorder, who are not eligible for inclusion. 6-Inability to communicate.

##### Intervention groups

For the intervention group, which consists of veterans with psychological injuries diagnosed with PTSD, the adaptive disclosure technique will be implemented in six weekly sessions, each lasting one and a half hours. The details of each session are as follows: Session 1: Assessment of the patient's current symptoms. Session 2: In-depth exploration of the patient's experiences related to the traumatic events experienced by veterans. Session 3: Addressing harmful thought patterns, developing effective coping strategies, and creating a trauma narrative. Session 4: Identifying core beliefs and challenges. Session 5: Setting new and positive goals, enhancing self-esteem, and reinforcing new behavioral

patterns. Session 6: Strengthening new behavioral patterns and assisting the patient in adapting to life after trauma.

##### Main outcome variables

severity of post-traumatic stress disorder symptoms

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20160815029377N4**

Registration date: **2025-03-05, 1403/12/15**

Registration timing: **registered\_while\_recruiting**

Last update: **2025-03-05, 1403/12/15**

Update count: **0**

##### Registration date

2025-03-05, 1403/12/15

##### Registrant information

##### Name

Mohammad Malekzadeh

##### Name of organization / entity

Yasuj University of Medical Sciences

##### Country

Iran (Islamic Republic of)

##### Phone

+98 74 3332 3379

##### Email address

mohamad.malekzadeh@yums.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2025-02-19, 1403/12/01

##### Expected recruitment end date

2025-03-20, 1403/12/30

**Actual recruitment start date**  
empty

**Actual recruitment end date**  
empty

**Trial completion date**  
empty

**Scientific title**  
Exploring the impact of the Adaptive Disclosure technique on reducing the severity of post-traumatic stress disorder symptoms in veterans: a randomized controlled trial

**Public title**  
Exploring the impact of the Adaptive Disclosure technique on reducing the severity of post-traumatic stress disorder symptoms in veterans: a randomized controlled trial

**Purpose**  
Education/Guidance

**Inclusion/Exclusion criteria**

**Inclusion criteria:**  
Age 18 years or older Psychiatric veterans with a diagnosis of post-traumatic stress disorder (co-occurring disorders such as depression, anxiety, or treated substance abuse or dependency problems are permitted.) Have sufficient insight to receive the adaptive disclosure technique through the Mental State Examination (MSE) Have informed consent to enter the study

**Exclusion criteria:**  
Lack of interest in continuing participation in the study. Suicidal behavior or homicide, requiring urgent or emergency evaluation or treatment within the past three months. Serious Axis I mental disorders such as psychotic disorders or Bipolar I disorder are not eligible. Simultaneous enrollment in any cognitive-behavioral therapy, group therapy, or any other treatment involving systematic disclosure of disturbing memories related to the establishment. Loss of self-insight. Inability to communicate.

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

**Gender**  
Both

**Phase**  
N/A

**Groups that have been masked**  
*No information*

**Sample size**  
Target sample size: **58**

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

**Randomization description**  
Randomized block assignment will be carried out using the online platform Sealedenvelope. For this purpose, considering the total sample size of 58 participants and the existence of two groups (intervention and control), the block sizes will be multiples of two (either 2 or 4). Therefore, based on the number of blocks, the resulting sequence, and the predefined code, participants will be

allocated to the two groups: intervention (A) and control (B).

**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

Research Ethics Committees of Yasuj University of Medical Sciences

##### Street address

Yasuj University of Medical Sciences, Research Deputy of Yasuj University of Medical Sciences

##### City

Yasuj

##### Province

Kohgiluyeh-va-Boyerahmad

##### Postal code

7591741417

#### Approval date

2023-07-09, 1402/04/18

#### Ethics committee reference number

IR.YUMS.REC.1402.063

## Health conditions studied

### 1

#### Description of health condition studied

Post-traumatic stress disorder

#### ICD-10 code

F43.1

#### ICD-10 code description

Post-traumatic stress disorder (PTSD)

## Primary outcomes

### 1

#### Description

قبل The severity of post-traumatic stress disorder (PTSD) symptoms is measured using the Post-Traumatic Stress Disorder Checklist (PCL), with a score range of 17 to 85. The score is obtained by summing the responses to 17 items based on a Likert scale. The cutoff score for diagnosing PTSD is set at 50 for military personnel samples and has been validated exclusively for military populations.

#### Timepoint

Before the intervention begins and immediately after the intervention ends

#### **Method of measurement**

Post-Traumatic Stress Disorder Checklist (PCL)

#### **Secondary outcomes**

empty

#### **Intervention groups**

##### **1**

#### **Description**

Intervention group: After obtaining the ethical approval code, clinical trial registration code, and other necessary permissions from Yasuj University of Medical Sciences, the researcher will visit Shahid Rajaee Hospital and Salman Hospital in Yasuj to collect data. Following coordination with hospital and department authorities, and providing adequate information regarding the nature of the research, as well as necessary explanations about the relevant questionnaire, the questionnaire will be distributed among veterans with post-traumatic stress disorder (PTSD) who meet the inclusion criteria and give informed consent. The completed questionnaires will be collected by the researcher. For the intervention group, the adaptive disclosure technique will be implemented in six weekly sessions, each lasting one and a half hours. In the first session, the patient's current symptoms will be assessed, a detailed history will be collected, and trauma exposure will be reviewed. In the second session, the patient's experiences with veterans' traumatic events will be explored in-depth, along with addressing "hot cognitions" that negatively affect emotions, interpersonal communication in decision-making, and problem-solving abilities. In the third session, harmful thought patterns will be addressed, effective coping strategies will be developed, and the trauma narrative will be constructed. In the fourth session, core beliefs and challenges will be identified, the impact of the trauma will be further explored, and healthier ways of thinking will be introduced. In the fifth session, new and positive goals will be set, self-esteem will be enhanced, and new behavioral patterns will be reinforced. The final session will focus on strengthening new behavioral patterns and assisting the patient in adapting to life after trauma. Through this session, the therapist will address any persistent symptoms or destructive behaviors while helping the patient develop a sense of self-efficacy and increase their ability to manage PTSD symptoms. The therapist will also guide the patient in anticipating and coping with potential difficulties, as well as provide resources for long-term support. During this period, no intervention will be provided to the control group. Immediately after the intervention, both groups will complete the research questionnaire again, and the collected data will be analyzed.

#### **Category**

Treatment - Other

##### **2**

#### **Description**

Control group: During the intervention period, no intervention will be provided to the control group.

#### **Category**

Treatment - Other

#### **Recruitment centers**

##### **1**

#### **Recruitment center**

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

Shahid Rajaee Hospital and Salman Hospital

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

Mohammad Malekzadeh

##### **Street address**

Kohgiluyeh and Boyer-Ahmad, Yasuj, Motahari Boulevard, Takhteh Street, below the Municipality Traffic Department.

##### **City**

Yasuj

##### **Province**

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##### **Postal code**

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##### **Phone**

+98 74 1333 1759

##### **Email**

mzh541@yahoo.com

#### **Sponsors / Funding sources**

##### **1**

#### **Sponsor**

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

Yasouj University of Medical Sciences

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

Seyed Amin Hossaini Motlagh

##### **Street address**

Yasuj, Mehran, Abu Zar Boulevard, Eram Boulevard, Research and Technology Department, Yasuj University of Medical Sciences.

##### **City**

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##### **Province**

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##### **Email**

aminhomo@yahoo.com

#### **Grant name**

#### **Grant code / Reference number**

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

Yes

#### **Title of funding source**

Yasouj 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**

Yasouj University of Medical Sciences

**Full name of responsible person**

Mohammad Malekzadeh

**Position**

Associate professor

**Latest degree**

Ph.D.

**Other areas of specialty/work**

Health Psychology

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**Latest degree**

Ph.D.

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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