

Clinical Trial Protocol

Iranian Registry of Clinical Trials

10 Jun 2026

Comparing the effectiveness of group-based acceptance and commitment therapy (ACT) and compassion-focused therapy (CFT) on psychological distress, work-family conflict, parenting stress, and quality of life in working mothers

Protocol summary

Study aim

The primary aim of this study is to compare the efficacy of ACT group Therapy with CFT group Therapy on psychological distress, work-family conflict, parenting stress, self-compassion, psychological flexibility, and quality of life among employed mothers.

Design

A single-blind, parallel-group, randomized controlled trial with paired randomization based on age (In order to control for the effects of age variation across the intervention groups), conducted on 56 participants.

Settings and conduct

The intervention sessions will be conducted on the Google Meet audio-visual online platform under the supervision of the supervisor at Taleghani Hospital, Tehran. The study population consists of employed mothers. Regarding blinding, a single-blind design will be used, meaning the individual responsible for data analysis will be unaware of which intervention each group received.

Participants/Inclusion and exclusion criteria

Inclusion criteria: Employed and married mothers in Tehran; at least one child under the age of 18; aged between 25 and 50; stable internet and devices such as a smartphone or computer; providing informed consent to participate in online sessions and to complete assessment tools. Exclusion criteria: Concurrent psychological or pharmacological treatment; substance or alcohol abuse.

Intervention groups

Intervention group 1: Ten 120-minute online sessions of ACT group therapy; Intervention group 2: Ten 120-minute online sessions of CFT group therapy.

Main outcome variables

Psychological Distress; Work-Family Conflict; Parenting Stress; Self-compassion; Psychological flexibility; Quality

of Life

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20260106068572N1**

Registration date: **2026-01-27, 1404/11/07**

Registration timing: **prospective**

Last update: **2026-01-27, 1404/11/07**

Update count: **0**

Registration date

2026-01-27, 1404/11/07

Registrant information

Name

Negin Sayqalzanan Mashhadi

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

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Recruitment status

recruiting

Funding source

Expected recruitment start date

2026-04-19, 1405/01/30

Expected recruitment end date

2026-07-21, 1405/04/30

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Comparing the effectiveness of group-based acceptance and commitment therapy (ACT) and compassion-focused therapy (CFT) on psychological distress, work-family conflict, parenting stress, and quality of life in working mothers

Public title

Comparison of the effectiveness of Acceptance and Commitment group therapy (ACT) and Compassion-Focused group therapy (CFT) on the psychological well-being of employed mothers

Purpose

Supportive

Inclusion/Exclusion criteria**Inclusion criteria:**

Married mothers with at least one child under the age of 18 Currently employed Access to stable internet and suitable equipment (such as a smartphone or computer) and commitment to regular attendance in online group therapy sessions Informed consent to participate in online therapy sessions and complete assessment tools at the pre-test, post-test, and follow-up stages

Exclusion criteria:

Concurrent receipt of psychological therapy or pharmacotherapy (or at least three months without a change in medication dosage). Substance or alcohol use. Work schedules or personal commitments that conflict with the individual's weekly online group therapy session times.

Age

From **25 years** old to **50 years** old

Gender

Female

Phase

N/A

Groups that have been masked

- Data analyster

Sample size

Target sample size: **56**

Randomization (investigator's opinion)

Randomized

Randomization description

- Phase 1 (Targeted Sampling): Employed mothers are selected based on specific criteria through online announcements (such as social media, relevant working mothers' groups, or online psychology clinics) or via local organizations and companies in Tehran. - Phase 2 (Paired Block Randomization): A finalized list of 56 selected participants is compiled and sorted in descending order of age. They are then divided into 28 paired blocks based on age proximity (the two oldest participants form the first pair, followed by the next two, and so on, until the final pair with the youngest participants). Within each paired block, one participant is randomly assigned to the Acceptance and Commitment Therapy (ACT) intervention

group and the other to the Compassion-Focused Therapy (CFT) intervention group. This method ensures statistical balance between the two groups regarding age and controls for the confounding effects of the age variable.

Blinding (investigator's opinion)

Single blinded

Blinding description

This study will utilize a single-blind design, in which the data analyst will be blinded to the type of intervention received by the participants. The final data obtained from the two different intervention groups will be provided to the statistical analyst as numeric codes, and they will not be informed which code corresponds to which therapy group.

Placebo

Not used

Assignment

Parallel

Other design features**Secondary Ids**

empty

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

The Ethics Committee of Shahid Beheshti University of Medical Sciences

Street address

13th Floor, Block A, Headquarters of the Ministry of Health, Treatment and Medical Education, Iran Sima Street, between Falamak South and Zarfashan Streets, Ghods (West) Town, Tehran

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1467664961

Approval date

2026-01-05, 1404/10/15

Ethics committee reference number

IR.SBMU.MSP.REC.1404.671

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

Mothers Employed Outside the Home

ICD-10 code**ICD-10 code description****Primary outcomes****1****Description**

Psychological Distress: individual's overall psychological state across three key domains—self-reported perceived stress, perceived anxiety, and perceived depression, in DASS-21 Scale

Timepoint

Prior to the commencement of group therapy interventions (pre-test), immediately following the completion of group therapy interventions (post-test), and four months after the conclusion of group therapy interventions (follow-up).

Method of measurement

The 21-item Depression, Anxiety and Stress Scale (DASS-21), developed by Lovibond and Lovibond (1995).

2

Description

Work-Family Conflict: A type of inter-role conflict in which the pressures arising from work and family domains are in some aspects incompatible or contradictory. Consequently, fulfilling one role (work or family) makes it difficult or problematic to fulfill the other role. This conflict can manifest as work-to-family interference (when work demands hinder the performance of family roles) or family-to-work interference (when family responsibilities disrupt work performance).

Timepoint

Prior to the commencement of group therapy interventions (pre-test), immediately following the completion of group therapy interventions (post-test), and four months after the conclusion of group therapy interventions (follow-up).

Method of measurement

The Work-Family Conflict Scale, developed by Carlson et al. in 2000, which consists of 18 items.

3

Description

Score in the Parenting Stress Index - Short Form (PSI-SF)

Timepoint

Prior to the commencement of group therapy interventions (pre-test), immediately following the completion of group therapy interventions (post-test), and four months after the conclusion of group therapy interventions (follow-up).

Method of measurement

The Parenting Stress Index-Short Form (PSI-SF) was developed by Abidin (1995). It is the abbreviated version of the original scale and consists of 36 items.

4

Description

Score in the Self-Compassion Scale - Short Form (SCS-SF)

Timepoint

Prior to the commencement of group therapy interventions (pre-test), immediately following the completion of group therapy interventions (post-test), and four months after the conclusion of group therapy interventions (follow-up).

Method of measurement

The Self-Compassion Scale (SCS) was initially developed

by Neff (2003) with 26 items. Subsequently, Raes et al. (2011) introduced its short-form version (SCS-SF) comprising 12 items.

5

Description

Psychological flexibility score based on the Acceptance and Action Questionnaire - Second Edition (AAQ-II).

Timepoint

Prior to the commencement of group therapy interventions (pre-test), immediately following the completion of group therapy interventions (post-test), and four months after the conclusion of group therapy interventions (follow-up).

Method of measurement

The Acceptance and Action Questionnaire (AAQ-II) was developed by Bond et al. to measure psychological flexibility and experiential avoidance. This questionnaire consists of 10 items.

6

Description

Quality of life score based on the World Health Organization Quality of Life questionnaire - Brief Version (WHOQOL-BREF).

Timepoint

Prior to the commencement of group therapy interventions (pre-test), immediately following the completion of group therapy interventions (post-test), and four months after the conclusion of group therapy interventions (follow-up).

Method of measurement

The World Health Organization Quality of Life-BREF questionnaire (WHOQOL-BREF)

Secondary outcomes

empty

Intervention groups

1

Description

First Intervention Group: The ACT group therapy will be conducted in 10 sessions, each lasting 120 minutes, based on the protocol published by Dara Wistrup and JoAnn Wright in 2017. ACT group therapy is a psychological intervention conducted in a group format, grounded in the principles of ACT. This approach, by focusing on accepting negative thoughts and emotions rather than combating them, encourages individuals to align their actions with their personal values and to engage in committed behaviors toward value-driven goals. Within this group therapy model, participants learn skills such as cognitive defusion, acceptance, and committed action through interpersonal interactions and mindfulness-based exercises, leading to enhanced psychological flexibility. Emphasizing six core processes (acceptance, cognitive defusion, being present in the moment, self-as-context, values, and committed action),

this approach aims to improve quality of life and reduce the negative impacts of psychological distress.

Category

Treatment - Other

2

Description

Second Intervention Group: The CFT group therapy will be conducted in 10 sessions, each lasting 120 minutes, based on the protocol published by Nicola Petrocchi and colleagues in 2024. CFT group therapy is a therapeutic group approach designed based on the compassion-focused therapy theory developed by Paul Gilbert. This method aims to cultivate compassion towards oneself and others, helping individuals manage difficult emotions such as shame, self-criticism, and anxiety using compassion-based techniques. In this form of group therapy, participants enhance their emotional regulation and self-acceptance skills through mindfulness exercises, guided imagery, and group discussions, which leads to reduced psychological distress and improved mental health. Utilizing the tripartite model of emotion regulation (threat system, drive system, and soothing system), this approach assists individuals in experiencing greater psychological balance.

Category

Treatment - Other

Recruitment centers

1

Recruitment center

Name of recruitment center

Tehran city

Full name of responsible person

Negin Sayqalzanan Mashhadi

Street address

Tehran - Shahid Chamran Highway - Yemen Street - Shahid A'rabi Street, adjacent to (next to) Ayatollah Taleghani Hospital.

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Shahid Beheshti University of Medical Sciences

Full name of responsible person

Hoda Doosalivand

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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

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

Negin Sayqalzanan Mashhadi

Position

PhD Assistant in Clinical Psychology

Latest degree

Ph.D.

Other areas of specialty/work

Psychology

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Latest degree

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Other areas of specialty/work

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Person responsible for updating data

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Name of organization / entity

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Full name of responsible person

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PhD Assistant in Clinical Psychology

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Sharing plan

Deidentified Individual Participant Data Set (IPD)

Yes - There is 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

Yes - There is a plan to make this available

Informed Consent Form

Yes - There is 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

Not applicable

Title and more details about the data/document

Our research team is able to share the final research data files as well as the results of statistical analyses upon completion of all implementation and writing phases of the study. However, no personally identifiable information of participants will be published in any reports or datasets. Confidentiality will be strictly maintained, and research data will only be released using randomly assigned numerical codes for each participant.

When the data will become available and for how long

Six months after the completion of the clinical trial.

To whom data/document is available

Only researchers affiliated with academic or scientific institutions will have access to the data.

Under which criteria data/document could be used

Data access will be granted exclusively for ethical and permissible scientific research purposes.

From where data/document is obtainable

Negin Sayalzanan Mashhadi
sayqalzanannegin@gmail.com

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

The interested researcher must email the designated contact person. After the research team approves the request, verifies the applicant's identity as a formal researcher affiliated with a reputable university or research institution, and clarifies their scientific purpose for accessing the data and documents, the raw research data files will be made available to them.

Comments