

Clinical Trial Protocol

Iranian Registry of Clinical Trials

10 Jun 2026

The Efficacy of Cognitive Behavior Hypnotherapy for Body Dysmorphic Disorder

Protocol summary

Summary

This study examined the effect of cognitive behavioral hypnotherapy on body dysmorphic disorder and depression and disability of these patients. This study utilized a single subject design which includes three baseline phases, treatment phase lasting fifteen sessions and follow-up phase. The study site was the city of Sanandaj in spring 2015. Three patients with body dysmorphic disorder participated in this study by purposive available sampling. The Yale-Brown Obsessive Compulsive Scale modified for Body Dysmorphic Disorder (BDD-YBOCS), Millon Clinical Multiaxial Inventory-III (MCMI-III) and clinical interview were used as diagnostic tools. The BDD-YBOCS, Beck Depression Inventory (BDI-II) and Sheehan Disability Scale (SDS), were used every week in baseline phase and twice a week in treatment condition at beginning of each session.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2016050425838N2**

Registration date: **2016-09-04, 1395/06/14**

Registration timing: **retrospective**

Last update:

Update count: **0**

Registration date

2016-09-04, 1395/06/14

Registrant information

Name

Morteza Abarin

Name of organization / entity

University of Kurdistan

Country

Iran (Islamic Republic of)

Phone

+98 17 3223 0046

Email address

m.abbarin@hum.uok.ac.ir

Recruitment status

Recruitment complete

Funding source

Vice chancellor for research University of Kurdistan

Expected recruitment start date

2015-04-04, 1394/01/15

Expected recruitment end date

2015-06-05, 1394/03/15

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The Efficacy of Cognitive Behavior Hypnotherapy for Body Dysmorphic Disorder

Public title

The Efficacy of Cognitive Behavior Hypnotherapy for Body Dysmorphic Disorder

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria: The age range between 18 and 48 years; BDD duration of at least 6 months; Read and write ability; Get score 21 or higher on BDD-YBOCS; Having a diagnosis of body dysmorphic disorder in the Diagnostic Interview Exclusion criteria: Having serious thoughts about suicide based on scores of suicide item of Beck Depression Inventory ;Having simultaneous psychotherapy; Having simultaneous pharmacotherapy for psychological disorders; Having cosmetic surgery during treatment

Age

From **18 years** old to **45 years** old

Gender
Both

Phase
N/A

Groups that have been masked
No information

Sample size
Target sample size: **3**

Randomization (investigator's opinion)
N/A

Randomization description

Blinding (investigator's opinion)
Not blinded

Blinding description

Placebo
Not used

Assignment
Single

Other design features
the study was single subject design

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee
Ethics committee of University of Kurdistan

Street address
Sanandaj, Pasdaran street

City
Sanandaj

Postal code

Approval date
2015-03-07, 1393/12/16

Ethics committee reference number
1179836

Health conditions studied

1

Description of health condition studied
body dysmorphic disorder

ICD-10 code
F45.2

ICD-10 code description
The essential feature is a persistent preoccupation with the possibility of having one or more serious and progressive physical disorders. Patients manifest persistent somatic complaints or a persistent preoccupation with their physical appearance. Normal

Primary outcomes

1

Description
body dysmorphic disorder severity

Timepoint
three baseline spaced one week- every session during treatment- four month follow up

Method of measurement
Yale-Brown Obsessive Compulsive Scale modified for Body Dysmorphic Disorder

Secondary outcomes

1

Description
depression severity

Timepoint
three baseline spaced one week- every session during treatment (twice a week)- four month follow up

Method of measurement
beck depression inventory (DBI-II)

2

Description
disability severity

Timepoint
three baseline spaced one week- every session during treatment (twice a week)- four month follow up

Method of measurement
Sheehan Disability Scale

Intervention groups

1

Description
cognitive behavior hypnotherapy 15 ninety-minute sessions

Category
Treatment - Other

Recruitment centers

1

Recruitment center

Name of recruitment center
through announcements in Sanandaj

Full name of responsible person
morteza abbarin

Street address
gorgan- hesam street- hesam building- second floor

City
gorgan

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Vice chancellor for research university of kurdestan

Full name of responsible person

madi zamestani

Street address

Pasdaran street, University of Kurdistan

City

Sanandaj

Grant name**Grant code / Reference number****Is the source of funding the same sponsor organization/entity?**

Yes

Title of funding source

Vice chancellor for research university of kurdestan

Proportion provided by this source

100

Public or private sector

empty

Domestic or foreign origin

empty

Category of foreign source of funding

empty

Country of origin**Type of organization providing the funding**

empty

Person responsible for general inquiries**Contact****Name of organization / entity**

University of Kurdistan

Full name of responsible person

mahdi zamestani

Position

Faculty of Psychology, University of Kurdistan

Other areas of specialty/work**Street address**

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Web page address**Person responsible for scientific inquiries****Contact****Name of organization / entity**

university of kurdestan

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Position

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Web page address**Person responsible for updating data****Contact****Name of organization / entity****Full name of responsible person**

morteza abarin

Position

MA in Clinical Psychology- Principal investigator

Other areas of specialty/work**Street address**

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gorgan

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Fax**Email**

morteza.abbarin@gmail.com

Web page address**Sharing plan****Deidentified Individual Participant Data Set (IPD)**

empty

Study Protocol

empty

Statistical Analysis Plan

empty

Informed Consent Form

empty

Clinical Study Report

empty

Analytic Code

empty

Data Dictionary

empty