

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

09 Jun 2026

### The effect of Memantine on negative symptoms in patients with chronic schizophrenia: a double blind and placebo controlled trial

#### Protocol summary

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#### Summary

The objective of this randomized, double-blind, placebo controlled study is to test the hypothesis that the addition of Modafinil would improve psychopathology in subjects with schizophrenia treated with Risperidone. 40 patients with chronic DSM-IV-diagnosed schizophrenia will receive Risperidone (6 mg/day) combined with either placebo (N=20) or 200 mg/day of Modafinil (N=20) for 8 weeks. Efficacy will be defined as the change from baseline to endpoint in score on the Positive and Negative Syndrome Scale (PANSS). Side effects will be also evaluated using checklist and Extra-pyramidal Symptoms Rating Scale.

#### Recruitment status

**Recruitment complete**

#### Funding source

Tehran University of Medical Sciences

#### Expected recruitment start date

2010-05-14, 1389/02/24

#### Expected recruitment end date

2010-07-19, 1389/04/28

#### Actual recruitment start date

empty

#### Actual recruitment end date

empty

#### Trial completion date

empty

#### General information

#### Acronym

#### IRCT registration information

IRCT registration number: **IRCT138903131556N16**

Registration date: **2010-06-06, 1389/03/16**

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

Last update:

Update count: **0**

#### Registration date

2010-06-06, 1389/03/16

#### Registrant information

#### Name

Shahin Akhondzadeh

#### Name of organization / entity

Roozbeh Psychiatric Hospital, Tehran University of Medical Sciences

#### Country

Iran (Islamic Republic of)

#### Phone

+98 21 5541 2222

#### Email address

#### Scientific title

The effect of Memantine on negative symptoms in patients with chronic schizophrenia: a double blind and placebo controlled trial

#### Public title

Modafinil and treatment of schizophrenia

#### Purpose

Treatment

#### Inclusion/Exclusion criteria

Inclusion Criteria:1- Age between 18-55,2- Diagnosis of Schizophrenia based on DSM IV 3-chronic Schizophrenia,4- Minimum Score of 60 on Positive and Negative Scale- Exclusion Criteria:1- Substance dependence,2- IQ <70,3-any other mental disorder on axis I, 4-Any serious medical or neurological problem ,5- receiving oral antipsychotic medications during the last week or receiving any depot antipsychotic medication during the last month,6- receiving ECT during the last 14 days,7- receiving nitrite

#### Age

From **18 years** old to **50 years** old

#### Gender

Both

## Phase

2

## Groups that have been masked

No information

## Sample size

Target sample size: 40

## Randomization (investigator's opinion)

Randomized

## Randomization description

## Blinding (investigator's opinion)

Double blinded

## Blinding description

## Placebo

Used

## Assignment

Parallel

## Other design features

## Secondary ids

empty

## Ethics committees

### 1

#### Ethics committee

##### Name of ethics committee

Tehran University of Medical Sciences

##### Street address

Keshavarz Blvd

##### City

Tehran

##### Postal code

#### Approval date

2009-07-06, 1388/04/15

#### Ethics committee reference number

6157

## Health conditions studied

### 1

#### Description of health condition studied

schizophrenia

#### ICD-10 code

F20

#### ICD-10 code description

Schizophrenia

## Primary outcomes

### 1

#### Description

Severity of schizophrenia

#### Timepoint

Baseline and weeks 2-4-6-8 after beginning of treatment

#### Method of measurement

Positive and Negative Syndrome Scale (PANSS)

## Secondary outcomes

empty

## Intervention groups

### 1

#### Description

Tablet Risperidone (6 mg/day) combined with 5 to 20 mg/day of Tablets Modafinil as intervention group for 8 weeks

#### Category

Treatment - Drugs

### 2

#### Description

Tablets Risperidone (6 mg/day) combined with Tablets placebo as control group for 8 weeks

#### Category

Placebo

## Recruitment centers

### 1

#### Recruitment center

##### Name of recruitment center

Roozbeh Hospital

##### Full name of responsible person

Prof. Shahin Akhondzadeh

##### Street address

Roozbeh Hospital

##### City

Tehran

## Sponsors / Funding sources

### 1

#### Sponsor

##### Name of organization / entity

Tehran University of Medical Sciences

##### Full name of responsible person

Dr. Akbar Fotouhi

##### Street address

Keshavarz Blvd

##### City

Tehran

#### Grant name

#### Grant code / Reference number

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

Yes

#### Title of funding source

Tehran University of Medical Sciences

#### 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**

**Person responsible for scientific inquiries**

**Contact**

**Name of organization / entity**

Tehran University of Medical Sciences

**Full name of responsible person**

Prof. Shahin Akhondzadeh

**Position**

Prof. of Clinical Psychopharmacology

**Other areas of specialty/work**

**Street address**

Roazbeh Hospital-South Kargar street

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**Email**

s.akhond@sina.tums.ac.ir

**Web page address**

**Person responsible for updating data**

**Contact**

**Name of organization / entity**

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**Email**

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**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*