

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

06 Jul 2026

### **Spironolacton as adjunctive treatment of symptoms in patients with schizophrenia: a double blind and placebo controlled trial**

#### **Protocol summary**

##### **Study aim**

The objective of this randomized, double-blind, placebo controlled study is to test the hypothesis that the addition of spironolacton would improve psychopathology in subjects with schizophrenia

##### **Design**

randomized, double-blind, placebo controlled study  
method of randomization: Random permuted Block  
Sample size: 52 Trial phase: 3

##### **Settings and conduct**

The study will be conducted among patients attending Qods Psychiatric Hospital-Sanandaj. Patients with schizophrenia who have admitted in Qods Hospital will be included in the trial provided the inclusion criteria will be met,

##### **Participants/Inclusion and exclusion criteria**

Inclusion criteria:1-Diagnosis of Schizophrenia based on DSM-5 criteria; 2- Age between 18-55; 3-Chronic Schizophrenia- duration of the disorder more than 2 years; 4-Minimum score of 20 in negative sub score. 5- being stable on an antipsychotic for the last 2 months( 6 months for clozapin) Exclusion criteria:1-Any serious medical or neurological problem; 2- IQ less than 70; 3- Substance dependence during the last 6 months(except for nicotine and caffeine); 4-receiving ECT during the last 2 months ; 6-Acute or chronic systemic diseases; 7- History of neurosurgery; 8- History of head trauma 9- allergy to spironolactone 10-receiving mood stabilizers 11.gynecomastia

##### **Intervention groups**

Intervention group:regular antipsychotic combined with 100mg/day spironolacton as intervention group for 8 weeks Control group: regular antipsychotic plus placebo for 8 weeks

##### **Main outcome variables**

Severity of schizophrenia

#### **General information**

##### **Reason for update**

##### **Acronym**

##### **IRCT registration information**

IRCT registration number: **IRCT20091229002935N8**

Registration date: **2019-12-07, 1398/09/16**

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

Last update: **2019-12-07, 1398/09/16**

Update count: **0**

##### **Registration date**

2019-12-07, 1398/09/16

##### **Registrant information**

##### **Name**

Farzin Rezaei

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

Kurdistan University of Medical Sciences

##### **Country**

Iran (Islamic Republic of)

##### **Phone**

+98 87 1356 1822

##### **Email address**

farrezaei@muk.ac.ir

##### **Recruitment status**

**Recruitment complete**

##### **Funding source**

##### **Expected recruitment start date**

2019-11-21, 1398/08/30

##### **Expected recruitment end date**

2020-09-21, 1399/06/31

##### **Actual recruitment start date**

empty

##### **Actual recruitment end date**

empty

##### **Trial completion date**

empty

## Scientific title

Spironolacton as adjunctive treatment of symptoms in patients with schizophrenia: a double blind and placebo controlled trial

## Public title

Spironolacton as adjunctive treatment of negative symptoms in patients with schizophrenia

## Purpose

Treatment

## Inclusion/Exclusion criteria

### Inclusion criteria:

Diagnosis of Schizophrenia based on DSM-5 criteria Age between 18-55 years old Chronic Schizophrenia- duration of the disorder more than 2 years Minimum score of 20 in negative sub score being stable on antipsychotic for the last 2 months(6 months for clozapine)

### Exclusion criteria:

Any serious medical or neurological problem IQ less than 70. Substance dependence during the last 6 months(except for nicotine and caffeine) receiving ECT during the last 2 months Acute or chronic systemic diseases History of head trauma exacerbation of symptoms(developing the active phase) hypersensitivity to spironolacton post-psychotic depression gynecomastia

## Age

From **18 years** old to **55 years** old

## Gender

Both

## Phase

3

## Groups that have been masked

- Participant
- Care provider
- Investigator
- Outcome assessor

## Sample size

Target sample size: **52**

## Randomization (investigator's opinion)

Randomized

## Randomization description

Participants allocated to intervention or control group via Random permuted blocks ( each block has four cases): AABB, ABAB, BBAA, BABA A:Intervention Group B:Control Group

## Blinding (investigator's opinion)

Double blinded

## Blinding description

The participants, clinicians and outcome evaluators will be blind regarding grouping and receiving placebo

## Placebo

Used

## Assignment

Parallel

## Other design features

## Secondary Ids

empty

## Ethics committees

### 1

#### Ethics committee

##### Name of ethics committee

Ethics Committee of Kurdistan University of Medical Sciences

##### Street address

No 1, Pasdaran Blvd.Kurdistan University of Medical Sciences, Sanandaj

##### City

Sanandaj

##### Province

Kurdistan

##### Postal code

1344666177

#### Approval date

2019-09-30, 1398/07/08

#### Ethics committee reference number

IR.MUK.REC.1398.144

## Health conditions studied

### 1

#### Description of health condition studied

schizophrenia

#### ICD-10 code

F20.0

#### ICD-10 code description

Paranoid schizophrenia

## Primary outcomes

### 1

#### Description

Severity of schizophrenia

#### Timepoint

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

#### Method of measurement

by Positive and Negative Symptoms Scale

## Secondary outcomes

empty

## Intervention groups

### 1

#### Description

Intervention group: Antipsychotic medication combined with 100 mg/day spironolacton as intervention group for 8 weeks

#### Category

Treatment - Drugs

### 2

#### Description

Control group: An antipsychotic combined with placebo as control group for 8 weeks

**Category**

Placebo

**Recruitment centers**

1

**Recruitment center**

**Name of recruitment center**

Qhods hospital- Sanandaj

**Full name of responsible person**

Dr Farzin Rezaei

**Street address**

Pasdarán Blvd- Qhods hospital - Sanandaj

**City**

Sanandaj

**Province**

Kurdistan

**Postal code**

1344666177

**Phone**

+98 87 3366 0025

**Fax**

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**Email**

farrezaei@muk.ac.ir

**Sponsors / Funding sources**

1

**Sponsor**

**Name of organization / entity**

Sanandaj University of Medical Sciences

**Full name of responsible person**

Dr Ebrahim Ghaderi

**Street address**

Deputy of research, Kurdistan university of Medical Sciences, Pasdarán Blvd., Sanandaj

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**Phone**

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**Email**

ebrahimghaderi@yahoo.com

**Grant name**

**Grant code / Reference number**

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

Yes

**Title of funding source**

Sanandaj 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**

Sanandaj University of Medical Sciences

**Full name of responsible person**

Farzin Rezaei

**Position**

Associate professor of psychiatry

**Latest degree**

Specialist

**Other areas of specialty/work**

Psychiatrics

**Street address**

Qhods Hospital, Pasdarán street, Sanandaj, Iran

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**Email**

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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**

Yes - There is a plan to make this available

**Informed Consent Form**

Yes - There is a plan to make this available

**Clinical Study Report**

Yes - There is a plan to make this available

**Analytic Code**

No - There is not a plan to make this available

**Data Dictionary**

No - There is not a plan to make this available

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

All data will be distributed through final report

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

from 2021 to 2026

**To whom data/document is available**

academic researchers

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

by email

**From where data/document is obtainable**

Dr Farzin Rezaei

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

farrezaei@muk.ac.ir

**Comments**