

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

10 Jun 2026

### The effect of compound "Ma-ol-asal" on clinical manifestation of the adult asthma

#### Protocol summary

##### Study aim

evaluation of the effect of compound "Ma-ol-asal" on clinical manifestation of the adult asthma 18-60 years old

##### Design

After getting Ethics committee reference number and IRCT registration, patients who have mild to moderate asthma 18-60 years that have diagnosed by lung specialist and have inclusion criteria, divide to case and control group.

##### Settings and conduct

taking a history. all patients were carefully examined, completed a Asthma Control Test (ACT). The classic treatment of asthma was performed for both groups, which include the use of fluticasone spray 250/25 mcg1 puff every 12 hours. In addition to the aforementioned treatment, the experimental group also receive compound "Ma-ol-asal".

##### Participants/Inclusion and exclusion criteria

Inclusion criteria: patients with asthma 18-60 years old that is referred to the pulmonary clinic of the Loghman Hospital Exclusion criteria: under 18 and over 60 years old patients with severe asthma and require hospitalization patients with underlying diseases such as CF,COPD, heart failure, GERD, bronchiectasis, Pulmonary embolism and sarcoidosis Addicts to cigarettes BMI > 29 pregnant and lactating women

##### Intervention groups

Asthmatic patients who receive compound "Ma-ol-asal" and classic treatment. Asthmatic patients who receive placebo and classic treatment.

##### Main outcome variables

Shortness of breath Cough Wheez Spray intake Patient's function Asthma control rate

#### General information

##### Reason for update

##### Acronym

#### IRCT registration information

IRCT registration number: **IRCT20181115041664N1**

Registration date: **2020-05-10, 1399/02/21**

Registration timing: **retrospective**

Last update: **2020-05-10, 1399/02/21**

Update count: **0**

#### Registration date

2020-05-10, 1399/02/21

#### Registrant information

##### Name

Narges Kaveh

##### Name of organization / entity

##### Country

Iran (Islamic Republic of)

##### Phone

+98 21 8877 3521

##### Email address

nkaveh@sbmu.ac.ir

#### Recruitment status

**Recruitment complete**

#### Funding source

#### Expected recruitment start date

2018-09-23, 1397/07/01

#### Expected recruitment end date

2018-12-21, 1397/09/30

#### Actual recruitment start date

empty

#### Actual recruitment end date

empty

#### Trial completion date

empty

#### Scientific title

The effect of compound "Ma-ol-asal" on clinical manifestation of the adult asthma

#### Public title

The effect of compound "Ma-ol-asal" on treatment of the adult asthma

### **Purpose**

Treatment

### **Inclusion/Exclusion criteria**

#### **Inclusion criteria:**

mild to moderate asthma 18-60 years old Referral to the pulmonary clinic of Loghman Hospital, Tehran, Iran

#### **Exclusion criteria:**

under 18 and over 60 years old patients with severe asthma and require hospitalization patients with underlying diseases such as CF,COPD, heart failure, GERD, bronchiectasis, Pulmonary embolism and sarcoidosis Addicts to cigarettes BMI > 29 pregnant and lactating women

### **Age**

From **18 years** old to **60 years** old

### **Gender**

Both

### **Phase**

3

### **Groups that have been masked**

- Participant
- Investigator

### **Sample size**

Target sample size: **80**

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

Randomized

### **Randomization description**

Randomization: Block Randomization unit: Individual  
Randomization tool: Random numbers table  
How to build a sequence: In the random numbers table, the random number is selected from the table and is selected in rows or columns in the up and down and left or right directions of the next. Because amount of 4 blocks conditions for the two groups is six, the numbers are greater than 6 and the zero ignored and each digit identifies the block.

### **Blinding (investigator's opinion)**

Double blinded

### **Blinding description**

Patients enter the study after studying and completing the form of informed consent and knowledge that drugs and placebo are not known. The researcher does not know the drug cods for medication or placebo.

### **Placebo**

Used

### **Assignment**

Parallel

### **Other design features**

## **Secondary Ids**

empty

## **Ethics committees**

### **1**

#### **Ethics committee**

**Name of ethics committee**

Ethics committee of Shahid-Beheshti University of Medical Sciences

### **Street address**

The Office of Management Research, Shahid Beheshti University of Medical Sciences, next to Taleghani Hospital, Evin, Shahid Chamran Highway

### **City**

Tehran

### **Province**

Tehran

### **Postal code**

1985717443

### **Approval date**

2019-01-06, 1397/10/16

### **Ethics committee reference number**

IR.SBMU.RETECH.REC.1397.828

## **Health conditions studied**

### **1**

#### **Description of health condition studied**

Asthma

#### **ICD-10 code**

J45

#### **ICD-10 code description**

Asthma

## **Primary outcomes**

### **1**

#### **Description**

Breathlessness

#### **Timepoint**

Before and after the intervention

#### **Method of measurement**

Asthma control Test

### **2**

#### **Description**

Cough

#### **Timepoint**

Before and after the intervention

#### **Method of measurement**

Asthma control Test

### **3**

#### **Description**

Wheez

#### **Timepoint**

Before and after the intervention

#### **Method of measurement**

Asthma control Test

### **4**

#### **Description**

Spray Intake

#### **Timepoint**

Before and after the intervention  
**Method of measurement**  
Asthma control Test

## 5

### **Description**

Patient`s function

### **Timepoint**

Before and after the intervention

### **Method of measurement**

Asthma control Test

## 6

### **Description**

Asthma control rate

### **Timepoint**

Before and after the intervention

### **Method of measurement**

Asthma control Test

## 7

### **Description**

B.M.I

### **Timepoint**

Before and after the intervention

### **Method of measurement**

kg/m<sup>2</sup>

## **Secondary outcomes**

### 1

#### **Description**

-

#### **Timepoint**

-

#### **Method of measurement**

-

## **Intervention groups**

### 1

#### **Description**

Intervention group: receiver 10cc syrup in 100cc lukewarm water, 3 times per day with Seretide spray 1 puff every 12 hours for 3 months. Each 100cc of compound honey syrup contains 2 gr. of Cinnamomum verum, 2 gr. of Elettaria cardamomum, 1 gr. Zingiber officinalRoscoe, 1 gr. Alpinia galangal and 1 gr. Crocus sativus with water and honey. Medicine is provided by Niac company.

#### **Category**

Treatment - Drugs

### 2

#### **Description**

Control group: Receiver 10cc placebo in 100cc lukewarm

water, 3 times per day with Seretide spray 1 puff every 12 hours for 3 months. Each 100cc of placebo syrup contains 0.1% sodium benzoate, 0.1% saccharin and 0.25% C.M.G. Placebo is provided by Niac company.

#### **Category**

Placebo

## **Recruitment centers**

### 1

#### **Recruitment center**

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

Loghman hospital

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

Narges Kaveh

##### **Street address**

Shams Alley, Valiasr Avenue

##### **City**

Tehran

##### **Province**

Tehran

##### **Postal code**

1985717443

##### **Phone**

+98 21 8877 3521

##### **Email**

sitm@sbtmu.ac.ir

## **Sponsors / Funding sources**

### 1

#### **Sponsor**

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

Shahid Beheshti University of Medical Sciences

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

Rasool Chooapani

##### **Street address**

Shahid Beheshti University of Medical Sciences, Next to Taleghani Hospital, Evin, Shahid Chamran Highway, Shahid Beheshti University of Medical Sciences

##### **City**

Tehran

##### **Province**

Tehran

##### **Postal code**

1985717443

##### **Phone**

+98 21 8877 3521

##### **Email**

sitm@sbtmu.ac.ir

#### **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**

Narges Kaveh

**Position**

PhD candidate

**Latest degree**

Medical doctor

**Other areas of specialty/work**

Traditional Medicine

**Street address**

No.8 Shams Alley, Vali-e-Asr Street

**City**

Tehran

**Province**

Tehran

**Postal code**

1985717443

**Phone**

+98 21 8877 3521

**Email**

arash27.nk@gmail.com

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

Shahid Beheshti University of Medical Sciences

**Full name of responsible person**

Rasool Choopani

**Position**

Assistant Professor

**Latest degree**

Ph.D.

**Other areas of specialty/work**

Traditional Medicine

**Street address**

No.8 Shams Alley, Vali-e-Asr Street

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**Email**

rchoopani@yahoo.com

**Person responsible for updating data****Contact****Name of organization / entity**

Shahid Beheshti University of Medical Sciences

**Full name of responsible person**

Narges Kaveh

**Position**

PhD candidate

**Latest degree**

Medical doctor

**Other areas of specialty/work**

Traditional Medicine

**Street address**

No.8 Shams Alley, Vali-e-Asr Street

**City**

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**Province**

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**Postal code**

1985717443

**Phone**

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**Email**

arash27.nk@gmail.com

**Sharing plan****Deidentified Individual Participant Data Set (IPD)**

No - There is not a plan to make this available

**Justification/reason for indecision/not sharing IPD**

We have not got any plans for dispersion.

**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**

No - There is not a plan to make this available

**Data Dictionary**

No - There is not a plan to make this available