

Clinical Trial Protocol

Iranian Registry of Clinical Trials

07 Jul 2026

Evaluation of the effect of Iranian medicine product, Algro, on the symptoms of patients with covid-19 (Coronavirus 19 disease)

Protocol summary

Study aim

Assessment of the effect of Iranian medicine product, Algro, on the symptoms of patients with covid-19 (Coronavirus 19 disease)

Design

This clinical trial has control group, and with parallel groups, double blinded, randomized, phase 2 to 3 and is performed on 80 patients. Randomization method will be blocked randomization (Quaternary random blocks).

Settings and conduct

Patients in the drug group take the Iranian traditional drug for ten days and 3 times daily In addition to the usual medicines (According to the protocol of the Ministry of Health). Patient in control group use routine treatment and placebo. The study is performed in Baqiyatallah Hospital.

Participants/Inclusion and exclusion criteria

Inclusion criteria: Male or female patients 18 years old and older with COVID-19, positive coronavirus 19 PCR, Ground glass view at low-dose CT scan, Arterial oxygen saturation less than 93%, Consciously completed consent form completed by the patient or the patient's supervisor. Exclusion criteria: History of pulmonary malignancy, History of asthma or COPD, History of Disabling disease or malignancy, Liver or kidney disorders.

Intervention groups

Intervention groups will be two ones. Patients in the intervention group will receive the intervention drug in addition to the routine treatment, but in control group, they will receive placebo in addition to the routine treatment.

Main outcome variables

3D CT Scans, dyspnea, cough, arterial oxygen saturation, PCR, CBC diff

General information

Reason for update

Acronym

Partial Traditional Persian medicine corona study :PTPMCS

IRCT registration information

IRCT registration number: **IRCT20160131026298N3**
Registration date: **2020-04-30, 1399/02/11**
Registration timing: **prospective**

Last update: **2020-11-02, 1399/08/12**

Update count: **1**

Registration date

2020-04-30, 1399/02/11

Registrant information

Name

Ahmad Reza Sharifi Olounabadi

Name of organization / entity

Baqiyatallah University of Medical Science,
Department of Traditional Iranian Medicine

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2020-05-04, 1399/02/15

Expected recruitment end date

2020-05-19, 1399/02/30

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Evaluation of the effect of Iranian medicine product, Algo, on the symptoms of patients with covid-19 (Coronavirus 19 disease)

Public title

Evaluation of the effect of Iranian medicine product, Algo, on patients with coronavirus 19

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria:

Male or female patients 18 years old and older with COVID-19 positive coronavirus 19 PCR Ground glass view at low-dose CT scan Arterial oxygen saturation less than 93% Consciously completed consent form completed by the patient or the patient's supervisor

Exclusion criteria:

History of pulmonary malignancy History of asthma or COPD History of Disabling disease or malignancy Liver or kidney disorders

Age

From **18 years** old

Gender

Both

Phase

2-3

Groups that have been masked

- Participant
- Care provider

Sample size

Target sample size: **80**

Randomization (investigator's opinion)

Randomized

Randomization description

Randomization method will be blocked randomization (Quaternary random blocks). Randomization units are individuals. Eighty outpatients referred to the hospital emergency department are randomly assigned to one of the two intervention and control groups. This study is a double-blind method and patients in the intervention group will receive the intervention drug in addition to the routine treatment, but in control group, they will receive the placebo in addition to the routine treatment.

Blinding (investigator's opinion)

Double blinded

Blinding description

Blinding will be performed using a placebo in the control group.

Placebo

Used

Assignment

Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethics committee of Baqiatallah University of Medical Sciences

Street address

Shahid Nosrati Alley, Sheikh Baha'i South Street, Mulla Sadra Street, Vanak Square

City

Tehran

Province

Tehran

Postal code

1435916471

Approval date

2020-03-29, 1399/01/10

Ethics committee reference number

IR.BMSU.REC.1399.038

Health conditions studied

1

Description of health condition studied

COVID-19

ICD-10 code

U07.2

ICD-10 code description

U07.2 COVID-19, virus not identified, COVID-19

Primary outcomes

1

Description

Measurement of cough severity

Timepoint

Day 1 before and at the end of treatment

Method of measurement

Standard cough questionnaire

2

Description

severity of shortness of breath

Timepoint

Day 1 before and at the end of treatment

Method of measurement

Shortness Of Breath With Daily Activity (SOBDA) Questionnaire

3

Description

lung radiologic changes

Timepoint

At beginning and end of the study

Method of measurement

Chest CT scan

Secondary outcomes

empty

Intervention groups

1

Description

Intervention group: The medicinal product includes a capsule composed of two herbs (salix, lambs quarter). The amount of each of these substances in the capsule is 250 mg. The dosage of the product is three capsules a day, which is half an hour before each meal. This product is prepared by Armaghan Sabz Araz Teb Pharmaceutical Company. In this group, common medical drug is also consumed according to the latest protocol of the Ministry of Health.

Category

Treatment - Drugs

2

Description

Control group: Patient in control group use common medicine based on the latest protocol of the Ministry of Health and placebo.

Category

Placebo

Recruitment centers

1

Recruitment center

Name of recruitment center

Baqiyatallah Hospital

Full name of responsible person

REZAMOHTASHAMI; AHMAD REZA SHARIFI
OLOUNABADI

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After Sheikh Bahaei avenue, Mulla Sadra Street,
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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Bagheiat-allah University of Medical Sciences

Full name of responsible person

Gholamhosein Alishiri

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Baqiyatallah University, Third Floor, Vice-Chancellor
for Research and Technology, Sheikh Bahai St, Mulla
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Grant name

Grant code / Reference number

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

No

Title of funding source

Armaghan Sabz Araz Teb Company

Proportion provided by this source

30

Public or private sector

Private

Domestic or foreign origin

Domestic

Category of foreign source of funding

empty

Country of origin

Type of organization providing the funding

Other

Person responsible for general inquiries

Contact

Name of organization / entity

Bagheiat-allah University of Medical Sciences

Full name of responsible person

Ahmad Reza Sharifi Olounabadi

Position

Assistant Professor

Latest degree

Ph.D.

Other areas of specialty/work

Traditional Medicine

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Person responsible for scientific inquiries

Contact

Name of organization / entity

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Full name of responsible person

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Position

Assistant Professor

Latest degree

Specialist

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Internal Medicine

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Latest degree

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Other areas of specialty/work

Traditional Medicine

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Sharing plan

Deidentified Individual Participant Data Set (IPD)

Undecided - It is not yet known if there will be 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

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

Undecided - It is not yet known if there will be a plan to make this available

Data Dictionary

Undecided - It is not yet known if there will be a plan to make this available