

Clinical Trial Protocol

Iranian Registry of Clinical Trials

08 Jun 2026

Evaluation of the effect of herbal food product (Shad syrup) in patients with COVID 19

Protocol summary

Study aim

Evaluation of the effect of herbal food product (Shad syrup) in patients with COVID19

Design

Clinical trial with control group, with parallel groups , non-blind, randomized using random numbers table, Phase three on 100 patients

Settings and conduct

Taleghani hospital, Tehran,Iran

Participants/Inclusion and exclusion criteria

inclusion criteria: confirmed COVID-19 infection with Lab tests not considering sign and symptoms suspected COVID-19; infection based on national protocol include the patients who has respiratory signs and pulmonary unilateral or bilateral multilobar infiltration in CT scan or chest radiography; suspected patient to pneumonia with uncommon rapid deterioration and fast worsening of clinical status and not responding to proper treatments; age 30 to 65. Exclusion criteria: Late visit more than 5 days of onset of symptoms; pregnancy and breastfeeding; allergy history; complicated bacterial infection severe and critical patients (ARDS and MODS); patient in recovery phase

Intervention groups

Intervention group: persacivir. Control group: placebo

Main outcome variables

death; fever (temperature); SPO2; CRP; ESR; LDH; SGPT; SGOT; BUN; Cr

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20150815023620N9**

Registration date: **2021-03-13, 1399/12/23**

Registration timing: **retrospective**

Last update: **2021-03-13, 1399/12/23**

Update count: **0**

Registration date

2021-03-13, 1399/12/23

Registrant information

Name

Mohammadreza Moshari

Name of organization / entity

Shahid Beheshti University of Medical Sciences

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2020-04-01, 1399/01/13

Expected recruitment end date

2020-08-01, 1399/05/11

Actual recruitment start date

2020-04-01, 1399/01/13

Actual recruitment end date

2020-08-01, 1399/05/11

Trial completion date

2020-08-01, 1399/05/11

Scientific title

Evaluation of the effect of herbal food product (Shad syrup) in patients with COVID 19

Public title

Evaluation of the effect of herbal food product in patients with COVID 19

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria:

confirmed COVID-19 infection with Lab tests not considering sign and symptoms suspected COVID-19 infection based on national protocol include the patients who has respiratory signs and pulmonary unilateral or bilateral multilobar infiltration in CT scan or chest radiography suspected patient to pneumonia with uncommon rapid deterioration and fast worsening of clinical status and not responding to proper treatments age 30 to 65

Exclusion criteria:

late visit more than 5 days of onset of symptoms pregnancy and breastfeeding allergy history complicated bacterial infection severe and critical patients (ARDS and MODS) patient in recovery phase

Age

From **30 years** old to **60 years** old

Gender

Both

Phase

3

Groups that have been masked

No information

Sample size

Target sample size: **150**

Actual sample size reached: **100**

Randomization (investigator's opinion)

Randomized

Randomization description

With simple randomization and using a random number table and individual randomization unit. To randomize, we use a table consisting of random digits 0 to 9. Each digit of this table is repeated the same on average. There is no pattern of recognizable numbers. In this method, each number is assigned to a treatment group. We start from the first line of the table and move down line by line. For the two treatments, we put the numbers 0 to 4 for treatment A and the numbers 5 to 9 for treatment B. The numbers in the first line of the table are as follows: 0,5,2,7,8,4,3,7,4,1,6,8,3,8,5,1,5,6,9,6, ... Now for people based on the above numbers, we have the following allocation: A, B, A, B, B,... We will continue the above process until the two groups are completed.

Blinding (investigator's opinion)

Not blinded

Blinding description**Placebo**

Used

Assignment

Parallel

Other design features**Secondary Ids**

empty

Ethics committees**1****Ethics committee**

Name of ethics committee

Vice-Chancellor in Research Affairs - Shahid Beheshti University of Medical Sciences

Street address

Anesthesiology Research Center,Taleghani Hospital Tehran, Iran

City

Tehran

Province

Tehran

Postal code

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Approval date

2020-05-30, 1399/03/10

Ethics committee reference number

IR.SBMU.RETECH.REC.1399.142

Health conditions studied**1****Description of health condition studied**

COVID-19

ICD-10 code

U07.1

ICD-10 code description

COVID-19, virus identified

Primary outcomes**1****Description**

Death

Timepoint

daily

Method of measurement

Patient record

2**Description**

Fever (temperature)

Timepoint

Daily

Method of measurement

Thermometer

3**Description**

SPO2

Timepoint

Daily

Method of measurement

Pulse oximetry

4**Description**

CRP

Timepoint

Daily

Method of measurement

Biochemical testing

5**Description**

ESR

Timepoint

Daily

Method of measurement

ESR Reader

6**Description**

LDH

Timepoint

Daily

Method of measurement

Biochemical testing

7**Description**

SGPT

Timepoint

Daily

Method of measurement

Biochemical testing

8**Description**

SGOT

Timepoint

Daily

Method of measurement

Biochemical testing

9**Description**

BUN

Timepoint

Daily

Method of measurement

Biochemical testing

10**Description**

Cr

Timepoint

Daily

Method of measurement

Biochemical testing

Secondary outcomes

empty

Intervention groups**1****Description**

Intervention group: Shad syrup. The dosage of the syrup is 5 cc with 10 cc of water every 4 hours for the first two days and 5 cc every 6 hours for the next two days, which is given to the patient by the ward nurse and its consumption is monitored.

Category

Treatment - Drugs

2**Description**

Control group: placebo. The dose is equivalent to non-drug syrup for the first two days every 4 hours 5 cc with 10 cc of water and the next two days every 6 hours 5 cc which is given to the patient by the ward nurse and its consumption is monitored.

Category

Placebo

Recruitment centers**1****Recruitment center****Name of recruitment center**

Tehran Taleghani hospital

Full name of responsible person

Reza Moshari

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Sponsors / Funding sources**1****Sponsor****Name of organization / entity**

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Full name of responsible person

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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
Sbmu Univercity
Proportion provided by this source
100
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
Academic

Person responsible for general inquiries

Contact
Name of organization / entity
Shahid Beheshti University of Medical Sciences
Full name of responsible person
Dr Mohammadreza Moshari
Position
Assistant professor
Latest degree
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Person responsible for updating data

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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

Yes - There is a plan to make this available

Data Dictionary

Yes - There is a plan to make this available

Title and more details about the data/document

some of information of main outcome will share

When the data will become available and for how long

from 2021

To whom data/document is available

researchers in universities

Under which criteria data/document could be used

use of all shared data is allowed with permission

From where data/document is obtainable

dr reza moshari rmoshari@sbmu.ac.ir

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

send an email and take a permission

Comments