

Clinical Trial Protocol

Iranian Registry of Clinical Trials

27 Jun 2026

Efficacy study of allogeneic human menstrual blood stem cells secretome to treat moderate to severe Covid-19 patients, clinical trial phase III

Protocol summary

derived allogeneic stem cells secretome in patients with moderate to severe pneumonia caused by Covid-19

Study aim

Evaluation of efficacy of injection of menstrual blood-derived stem cells secretome in patients with Covid-19 moderate to severe pneumonia

Design

A clinical trial with the control group, with parallel groups, double-blind, randomized, phase 3 on 90 patients. Web random rendering software will be used for randomization.

Settings and conduct

The study population is selected according to the inclusion and exclusion criteria and consent is obtained from all and then these people are divided into two groups of intervention and control groups using randomization. The culture medium of allogeneic menstrual blood stem cells is collected in a cleanroom of Ibn Sina Research Institute and evaluated for contamination and then injected intravenously into patients with severe pneumonia caused by Covid-19.

Participants/Inclusion and exclusion criteria

Age 20 to 75 years, confirmed pneumonia caused by Covid-19, a positive test (RT-PCR) for Covid-19, diagnosed with severe disease: shortness of breath and respiratory distress, Respiratory rate ≥ 30 times/min; % of blood oxygen saturation at rest $\leq 90\%$; PaO₂/FiO₂ ratio ≤ 300 mmHg; Pulmonary infiltration more than 50% within 24 to 48 hours or moderate disease: Presence of respiratory symptoms with or without fever equal to/greater than 38 ° C; Percentage of blood oxygen saturation in Room air at rest between 90 to 93%; Pulmonary infiltration less than 50%

Intervention groups

Intervention group: treat with the fusion of the allogeneic supernatant of menstrual blood stem cells. Control group: in addition to receiving routine national treatments, undergo intravenous injection of normal saline.

Main outcome variables

Determination of efficacy of injection of menstrual blood-

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20180619040147N7**

Registration date: **2021-08-12, 1400/05/21**

Registration timing: **registered_while_recruiting**

Last update: **2021-08-12, 1400/05/21**

Update count: **0**

Registration date

2021-08-12, 1400/05/21

Registrant information

Name

Maryam Darzi

Name of organization / entity

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2021-07-23, 1400/05/01

Expected recruitment end date

2021-10-22, 1400/07/30

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Efficacy study of allogeneic human menstrual blood stem cells secretome to treat moderate to severe Covid-19 patients, clinical trial phase III

Public title

Treatment of Covid-19 using menstrual blood stem cell secretome

Purpose

Treatment

Inclusion/Exclusion criteria**Inclusion criteria:**

Aged 40-65 years Voluntarily participate in this clinical trial and sign off "informed consent form" Chest imaging confirm COVID-19 featured lesions in the lung The SARS-CoV-2 nucleic acid test was positive Diagnosed with severe pneumonia of COVID-19: respiratory distress, Respiratory rate (RR) \geq 30 times/min; resting oxygen saturation of 90% or less; arterial pressure of oxygen/the fraction of inspired oxygen \leq 300 mmHg; pulmonary imaging of focus within 24-48 hours > 50% progression Patients with a diagnosis of moderate pneumonia caused by Covid-19: 1- Presence of respiratory symptoms (including shortness of breath, feeling of pain and pressure in the chest, ...) with or without fever equal to / greater than 38 ° C 2- Percentage of blood oxygen saturation in Room air at rest between 90 to 93% 3- Pulmonary infiltration less than 50%

Exclusion criteria:

History of drug reactions or allergies Pneumonia caused by bacteria, Mycoplasma, Chlamydia, Legionella, fungi, or other viruses Airway obstruction due to lung cancer or unknown factors Carcinoid syndrome History of epilepsy and long-term use of anticonvulsant drugs during the last 3 years History of long-term use of immunosuppressive drugs History of chronic respiratory illness that requires long-term oxygen therapy The patient is on blood or peritoneal dialysis Creatinine clearance <15 ml / min Moderate to severe liver disease (Child-Pugh score > 12) History of deep vein thrombosis (DVT) or pulmonary embolism over the past 3 years Being under ECMO or high-frequency oscillatory ventilation support Diagnostic of HIV, hepatitis B, and syphilis Pregnant or lactating women Lack of consciousness and inability to provide informed consent by the patient

Age

From **20 years** old to **75 years** old

Gender

Both

Phase

3

Groups that have been masked

- Participant
- Data analyser

Sample size

Target sample size: **90**

Randomization (investigator's opinion)

Randomized

Randomization description

In this study, a simple computer-aid randomization method is used. In this method, a list of numbers from 1 to 90 is prepared, and each number is randomly assigned to groups A or B by a computer. Depending on the time of hospitalization, these numbers are assigned to the patients, respectively, and based on the list of patients, they are assigned to intervention group A (routine treatment with cell therapy) and control group B (routine treatment with normal saline injection). So that in each group there are 45 patients.

Blinding (investigator's opinion)

Double blinded

Blinding description

This study is a double-blind trial (participants and data and outcome assessors). Because the drug is in the form of a stem cell supernatant cultured in a phenol-free culture medium, this medium is similar in color and volume to a normal saline injection serum, and therefore at the time of injection, The patient will not notice any difference in the color or volume of the medicine. The data analyst will also not know which of the drug/placebo options each patient receives and is unaware of the nature of the codes assigned to patients.

Placebo

Used

Assignment

Parallel

Other design features**Secondary Ids**

empty

Ethics committees**1****Ethics committee****Name of ethics committee**

Research Ethics Committees of Avicenna Research Institute

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Avicenna Research Institute, Shahid Beheshti University, Shahid Chamran Highway, Tehran, Iran

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Postal code

1983969412

Approval date

2021-07-19, 1400/04/28

Ethics committee reference number

IR.ACECR.AVICENNA.REC.1400.008

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

moderate to severe covid-19 patients

ICD-10 code

U07.1

ICD-10 code description

COVID-19, virus identified

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

Decrease in serum CRP levels

Timepoint

On days 0, 3, 5, 7, 10, 14, 21, 28 after the first intervention

Method of measurement

Laboratory evaluation

2**Description**

Decrease in serum levels of lactate dehydrogenase

Timepoint

On days 0, 3, 5, 7, 10, 14, 21, 28 after the first intervention

Method of measurement

Laboratory evaluation

3**Description**

Decrease in serum D-Dimer levels

Timepoint

On days 0, 3, 5, 7, 10, 14, 21, 28 after the first intervention

Method of measurement

Laboratory evaluation

4**Description**

Decrease in lymphopenia percentage

Timepoint

On days 0, 3, 5, 7, 10, 14, 21, 28 after the first intervention

Method of measurement

Laboratory evaluation

5**Description**

Reduce the size of the lesion on CT scan of the lungs

Timepoint

En On days 0, 10, 28 after the first intervention

Method of measurement

CT Scan

Secondary outcomes**1****Description**

Improving respiratory efficiency

Timepoint

Before the intervention until the one month after the first intervention on a daily basis

Method of measurement

Measuring PaO₂ / FiO₂ ratio or percentage of blood oxygenation

2**Description**

Increasing the number of patients weaning from mechanical ventilation

Timepoint

Daily for one month after the first injection

Method of measurement

Observation

3**Description**

Reduce the number of days hospitalized in the ICU

Timepoint

Daily for one month after the first injection

Method of measurement

Observation

4**Description**

Reducing the incidence of failure of various organs

Timepoint

Daily for one month after the first injection

Method of measurement

Observation

5**Description**

Reduce mortality rate

Timepoint

Daily for one month after the first injection

Method of measurement

Counting people

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

Intervention group: patients are treated by intravenous injection of 5 times supernatant culture medium of allogeneic menstrual blood-derived stem cells. After isolation and culture in a cleanroom under the GMP of Sina Eltiem Biomedical Engineering Company GMP and passing and quality control tests, stem cells are stored in the cell bank. After cell thawing and culture in a GMP-approved culture medium, this surface culture medium is collected and centrifuged to remove cells and cell debris, and after filtration, quality control tests are performed on the collected culture medium. The 5 ml cell secretion is then filled into sterile vials and packaged. Each vial dissolved in 100 ml of normal saline is injected through a peripheral vein over a period of 60 minutes.

Category

Treatment - Drugs

2**Description**

Control group: in addition to receiving routine national treatments for this disease, 45 patients undergo intravenous injection of 100 ml of normal saline 5 times in completely similar conditions to the intervention group

Category

Placebo

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

Imam Khomeini hospital

Full name of responsible person

Ali Dehghan Manshadi

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Dr. Gharib Street, At the end of Keshavarz Blvd, Imam Khomeini Hospital Complex

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

Iranian academic center for education culture and research

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Grant name**Grant code / Reference number****Is the source of funding the same sponsor****organization/entity?**

Yes

Title of funding source

Iranian academic center for education culture and research

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**

Tehran University of Medical Sciences

Full name of responsible person

علی دهقان منشادی

Position

Associate Professor

Latest degree

Specialist

Other areas of specialty/work

Infectious diseases

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

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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

No - There is not 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

Not applicable

Data Dictionary

Not applicable