

Clinical Trial Protocol

Iranian Registry of Clinical Trials

13 Sep 2026

The effect of Plasmapheresis on hypoxia in patients suffering from COVID-19: A randomized clinical trial

Protocol summary

Study aim

Determination of the effect of plasmapheresis on hypoxia and inflammatory factors in patients with new coronavirus (Covid-19).

Design

Clinical trial with control group, with parallel groups, open-label, randomized, phase 3 on 40 patients. Simple randomization method was used as coding method. (Odd and even)

Settings and conduct

Patients referred to Hazrat Vali-e-Asr Hospital in Birjand with Covid-19 infection whose disease has been confirmed by real-time PCR molecular test or on the basis of lung CT scan symptoms and their infection is severe or threatening. Life is randomly assigned to the case and control groups and the study was conducted as an open label. In the intervention group, in addition to the standard treatment of Covid-19, plasmapheresis was also performed for the patients.

Participants/Inclusion and exclusion criteria

Definitive diagnosis of Covid-19 based on taking a sample from the pharynx and laboratory PCR confirmation, lung involvement of more than 50% and SPO2 levels below 80% without oxygen or below 88% with oxygen or IL above 10 or ferritin above 1500, patients who have not responded to other Covid-19 treatments, age over 18 and consent to participate in the study; Exclusion criteria include a history of albumin sensitivity, a history of heparin allergy who cannot receive heparin as an anticoagulant during plasmapheresis, patients with hypocalcemia.

Intervention groups

Patients in the intervention group, in addition to the standard treatment of Covid-19, will undergo the intervention (plasmapheresis). For each patient, if the clinical symptoms do not improve, plasmapheresis will be performed up to 3 times. In this group, patients received the standard treatment of Covid-19 according to the national protocol.

Main outcome variables

Percentage of blood oxygen saturation, Arterial blood oxygen pressure in the analysis of blood gases

General information

Reason for update

Acronym

BPS

IRCT registration information

IRCT registration number: **IRCT20220517054899N1**

Registration date: **2022-08-13, 1401/05/22**

Registration timing: **registered_while_recruiting**

Last update: **2022-08-13, 1401/05/22**

Update count: **0**

Registration date

2022-08-13, 1401/05/22

Registrant information

Name

Vajihollah Raeesi

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 56 3239 5000

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2022-07-22, 1401/04/31

Expected recruitment end date

2023-03-22, 1402/01/02

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The effect of Plasmapheresis on hypoxia in patients suffering from COVID-19: A randomized clinical trial

Public title

The effect of Plasmapheresis on hypoxia in patients suffering from COVID-19

Purpose

Treatment

Inclusion/Exclusion criteria**Inclusion criteria:**

Definitive diagnosis of Covid-19 based on taking a sample from the pharynx and PCR laboratory confirmation Lung involvement more than 50% SPO2 levels below 80% without oxygen or below 88% with oxygen or IL above 10 or ferritin above 1500 Patients who have not responded to other Covid-19 treatments Age over 18 years and consent to participate in the study

Exclusion criteria:

History of albumin sensitivity Heparin allergy who cannot receive heparin as an anticoagulant during plasmapheresis Patients with hypocalcemia (citrate used in the plasmapheresis process exacerbates hypocalcemia)

AgeFrom **18 years** old**Gender**

Both

Phase

3

Groups that have been masked*No information***Sample size**Target sample size: **40****Randomization (investigator's opinion)**

Randomized

Randomization description

Simple randomization based on the table of random numbers. A set of numbers without a specific pattern and order and in a completely random manner are in the form of a table. The direction of reading the numbers in this table will be from top to bottom and right to left. Even numbers will be considered for the intervention group and odd numbers for the control group.

Blinding (investigator's opinion)

Not blinded

Blinding description**Placebo**

Not used

Assignment

Parallel

Other design features**Secondary Ids**

empty

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

Ethics committee of Birjand University of Medical Sciences

Street address

No. 28, Ghafari Ave., zakariyay razi Blvd ,birjand University of Medical Sciences

City

birjand

Province

South Khorasan

Postal code

9717964151

Approval date

2022-04-27, 1401/02/07

Ethics committee reference number

IR.BUMS.REC.1401.059

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

Corona virus (Covid-19)

ICD-10 code

Pneumonia

ICD-10 code description

J12.81

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

Percentage of blood oxygen saturation (SPO2)

Timepoint

On days zero (before the intervention), days one, three, five and seven after the intervention

Method of measurement

Pulse oximeter Bitmos GmbH made in Germany

2**Description**

Arterial blood oxygen pressure in the analysis of blood gases Pao2

Timepoint

On days zero (before the intervention), days one, three, five and seven after the intervention

Method of measurement

Blood sample to measure blood gases

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

Duration of hospitalization of the patient
Timepoint
End of hospitalization or death of the patient
Method of measurement
By day

2

Description

C-reactive protein (CRP) Factor

Timepoint

On days zero (before the intervention), days one, three, five and seven after the intervention

Method of measurement

Blood Test Prestige 24i machine made in Japan

Intervention groups

1

Description

Intervention group: Patients in the intervention group, in addition to the standard treatment of Covid-19, (dexamethasone at a dose of 6 mg per day and remdesivir at a dose of 200 mg on the first day and then 100 mg daily for 7 days) will undergo the intervention (plasmapheresis). Each time plasmapheresis is performed, 40 ml per kg of body weight of the patient's plasma is replaced with 5% human albumin and 0.9% normal saline, and for each patient, if the clinical symptoms do not improve, plasmapheresis will be performed up to 3 times.

Category

Treatment - Drugs

2

Description

Control group: In this group, patients received the standard treatment of Covid-19 according to the national protocol (Hedhexamethasone 6 mg daily and Ramdesivir 200 mg on the first day and 100 mg daily for 7 days).

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Valiasr Hospital

Full name of responsible person

Vajehallah Raeesi

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Birjand University of Medical Sciences

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

Birjand 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

Birjand University of Medical Sciences

Full name of responsible person

Vajehallah Raeesi

Position

Assistant Professor

Latest degree

Subspecialist

Other areas of specialty/work

Urology

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Full name of responsible person

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Position

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Latest degree

Subspecialist

Other areas of specialty/work

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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

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

Title and more details about the data/document

-

When the data will become available and for how long

-

To whom data/document is available

-

Under which criteria data/document could be used

-

From where data/document is obtainable

-

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

-

Comments

-