

Clinical Trial Protocol

Iranian Registry of Clinical Trials

27 Jun 2026

Evaluation of the effect of pentoxifylline in reducing the complications of Covid_19

Protocol summary

Study aim

Evaluation of the effect of pentoxifylline in reducing the complications of Covid_19

Design

Randomized, non-blind, clinical trial with parallel intervention and control group, with 100 Covid 19 patients.

Settings and conduct

This study was done on 100 Covid_19 patients in Sina hospital of Tabriz University of Medical Sciences. Patients will receive pantoxifylline 400 mg tablet thrice daily with meal.

Participants/Inclusion and exclusion criteria

Inclusion criteria are: PCR Positive from nasopharyngeal specimen for covid 19 or evidence indicating Covid 19 disease, Moderate to mild symptoms, Less than 7 days after the onset of symptoms Ability to swallow, Age between 18 and 75 and Exclusion criteria are: Shock or failure of several organs, Chronic liver disease, Chronic kidney disease, Pregnancy and lactation, Participation in other clinical studies and Concomitant use of drugs that interact severely with pentoxifylline.

Intervention groups

This randomized, non blind clinical study will be performed on 100 selected patients according to the inclusion and exclusion criteria. Intervention group will receive 400 mg pentoxifylline tablets 3 times a day for 1 month with the standard diet of the Ministry of Health. The condition of patients will be evaluated daily with laboratory and clinical parameters and finally recorded in predesignated checklists. The control group will receive only the standard regimen of the Ministry of Health for 1 month.

Main outcome variables

Serum Lactate Dehydrogenase (LDH) level , Serum C_reactive protein (CRP) level , Percentage of Oxygen saturation in arterial blood (SPO2), prothrombin time (PT) and Partial Thromboplastin Time (PTT) , Serum Procalcitonin level , Serum Ferritin and D_dimer level and

Serum level Brain natriuretic peptide (NT_ProBNP).

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20170609034406N8**

Registration date: **2021-08-26, 1400/06/04**

Registration timing: **prospective**

Last update: **2021-08-26, 1400/06/04**

Update count: **0**

Registration date

2021-08-26, 1400/06/04

Registrant information

Name

Afshin Gharekhani

Name of organization / entity

Faculty of Pharmacy/Tabriz University of Medical Sciences

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2021-09-23, 1400/07/01

Expected recruitment end date

2022-02-19, 1400/11/30

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date
empty

Scientific title
Evaluation of the effect of pentoxifylline in reducing the complications of Covid_19

Public title
Investigation of the effect of pentoxifylline on Covid-19

Purpose
Treatment

Inclusion/Exclusion criteria
Inclusion criteria:
Positive PCR test for nasopharyngeal sample or certain evidence indicating Covid-19 disease Moderate to mild symptoms onset of symptoms within 7 days Ability to swallow Age between 18 and 75
Exclusion criteria:
Shock state or multi-organ failure Chronic liver disease Chronic kidney disease Pregnancy and lactation Participation in other clinical studies Receiving drugs with severe interaction with pentoxifylline

Age
From **18 years** old to **75 years** old

Gender
Both

Phase
3

Groups that have been masked
No information

Sample size
Target sample size: **100**

Randomization (investigator's opinion)
Randomized

Randomization description
In this study, Permuted Block Randomization method will be used for entering patients into control and treatment groups. There will exist 25 blocks with equal number of patients from each group in this study. Random numbers will be generated by using the Microsoft Excel Spreadsheet Software to randomize blocks and patients allocation.

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 Tabriz University of Medical Sciences

Street address

Research Vice-Chancellor, Third floor, No 2 central building, Tabriz University of Medical Sciences, Golgasht Street, Tabriz

City

Tabriz

Province

East Azarbaijan

Postal code

5766414766

Approval date

2021-02-14, 1399/11/26

Ethics committee reference number

IR.TBZMED.REC.1399.1045

Health conditions studied

1

Description of health condition studied

COVID_19

ICD-10 code

U10.9

ICD-10 code description

Multisystem inflammatory syndrome associated with COVID_19

Primary outcomes

1

Description

Serum Procalcitonin level

Timepoint

At the beginning and end of the study

Method of measurement

Procalcitonin kit

2

Description

Serum Ferritin level

Timepoint

At the beginning and end of the study

Method of measurement

Ferritin kit

3

Description

Serum NT_ProBNP level (Brain natriuretic peptide)

Timepoint

At the beginning and end of the study

Method of measurement

Nt_ProBNP ELISA Kit

4

Description

Serum D_dimer level

Timepoint

At the beginning and end of the study.

Method of measurement

D_dimer ELISA kit

5

Description

Serum LDH (Lactate Dehydrogenase) level

Timepoint

At the beginning and end of the study

Method of measurement

DGKC Kit

6

Description

Partial Thromboplastin Time (PTT) and prothrombin Time (PT)

Timepoint

At the beginning and end of the study

Method of measurement

PTT and PT Kit

7

Description

Saturation of Peripheral Oxygen in arterial blood (SPO2)

Timepoint

At the beginning and end of the study.

Method of measurement

Arterial Blood Gas (ABG Test)

8

Description

Serum C_ reactive protein (CRP) level

Timepoint

At the beginning and end of the study

Method of measurement

CRP ELISA Kit

Secondary outcomes

1

Description

Reducing the length of hospital stay of patients

Timepoint

At the end of study

Method of measurement

Comparing the number of hospitalization days between the intervention group and control group

2

Description

Improving the quality of lung lesions

Timepoint

At the beginning and end of the study.

Method of measurement

CT_Scan of lungs

3

Description

Change in the sense of smell

Timepoint

At the beginning and end of the study.

Method of measurement

Questionnaire

Intervention groups

1

Description

Intervention group (Pentoxifylline recipient + standard care): 100 patients with COVID-19 diagnosis will be included in the study according to the inclusion and exclusion criteria and will receive 400mg pentoxifylline tablet (produced by Amin pharmaceutical company in Iran) three times daily for 1 month along with the standard care of the Ministry of Health protocol.

Category

Treatment - Drugs

2

Description

Control group: In this clinical study, the control group will receive only the standard care of the Ministry of Health protocol for COVID-19 patients for 1 month.

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Sina Hospital of Tabriz University of Medical Sciences

Full name of responsible person

Dr Afshin Gharekhani

Street address

No 2 central building, Tabriz University of Medical sciences, Golgasht Street, Tabriz

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Tabriz 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?**

Yes

Title of funding source

Tabriz 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**

Tabriz University of Medical Sciences

Full name of responsible person

Dr Afshin Gharekhani

Position

University faculty member

Latest degree

Ph.D.

Other areas of specialty/work

Medical Pharmacy

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Person responsible for scientific inquiries

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Full name of responsible person

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Position

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Latest degree

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Other areas of specialty/work

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Person responsible for updating data

Contact**Name of organization / entity**

Tabriz University of Medical Sciences

Full name of responsible person

Atena Hatami Fard

Position

Pharmacy student at Tabriz University of Medical Sciences

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Bachelor

Other areas of specialty/work

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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
There is no further information.
Study Protocol
Yes - There is 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
Yes - There is a plan to make this available
Clinical Study Report
Yes - There is 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

Demographic data and main outcomes will be shared.
When the data will become available and for how long
Data files will become available 6 months after publication.
To whom data/document is available
It will be only available for people working in academic institutions.
Under which criteria data/document could be used
All of the data can be freely used if the citation is appropriately considered .
From where data/document is obtainable
The applicants will be referred to research Vice-chancellor .
What processes are involved for a request to access data/document
All of the requested data should be mentioned in a application letter which will be sent to the Research Vice-chancellor of Tabriz University of Medical Sciences.
Comments