

Clinical Trial Protocol

Iranian Registry of Clinical Trials

13 Sep 2026

Comparing the efficacy of Molnupiravir and placebo on recovery rate in patients with mild COVID-19, a randomized multicenter clinical trial

Protocol summary

Study aim

Evaluation of the efficacy of Molnupiravir in recovery pace of patients with mild COVID-19

Design

Phase 2 and 3 double-blind clinical trials with parallel groups that will be randomized using quadruple blocks and will be performed on 500 patients in two intervention groups with Molnupiravir and placebo groups

Settings and conduct

This research project will be performed in patients with COVID-19 in the initial phase who have referred to the Infectious Diseases Clinic of Baqiyatallah Hospital in Tehran. The study will be double-blinded and patients and outcome assessors will be blinded to the intervention groups. Drugs and placebo will be exactly the same, and drug labels will remain unknown to researchers until the end of the study.

Participants/Inclusion and exclusion criteria

Inclusion criteria: having a patient in the family; Positive COVID-19 PCR test; Onset of clinical symptoms of the patient less than 3 days; Willing and able to take oral medication; Blood oxygen level above 93%. Exclusion criteria: pregnancy; Breastfeeding; any underlying diseases; history of allergy to antiviral drugs; thrombocytopenia; receiving recent vaccination of COVID-19

Intervention groups

Intervention group: Molnupiravir 200 mg daily for up to five days with diphenhydramine syrup / vitamin C tablets / vitamin D tablets / famotidine tablets 40 mg every 12 hours and naproxen in case of muscle pain and fever. Control group: similar intervention with the difference that they will receive placebo instead of molnupiravir.

Main outcome variables

PCR test and CT number; Fever; dyspnea; Cough; respiratory rate; Serum ESR; Serum CRP; pulse rate; blood pressure; Dry cough

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20210914052480N2**

Registration date: **2022-07-07, 1401/04/16**

Registration timing: **registered_while_recruiting**

Last update: **2022-07-07, 1401/04/16**

Update count: **0**

Registration date

2022-07-07, 1401/04/16

Registrant information

Name

Alireza Soleymanitabar

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 21 8755 4364

Email address

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2022-04-21, 1401/02/01

Expected recruitment end date

2023-03-20, 1401/12/29

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Comparing the efficacy of Molnupiravir and placebo on recovery rate in patients with mild COVID-19, a randomized multicenter clinical trial		Assignment Parallel
Public title The efficacy of molnupiravir in patients with COVID-19		Other design features
Purpose Treatment		Secondary Ids empty
Inclusion/Exclusion criteria Inclusion criteria: Living in a family with at least one infected person with COVID-19 Not having pulmonary involvement Not having symptoms like chest wall pain and dysnea suggesting pulmonary involvement Positive PCR test of COVID-19 Patient with COVID-19 symptoms lesser than 3 days Patients satisfied with consumption of oral medication Blood oxygen saturation above 93 percents Exclusion criteria: Patients under 18 years old Having underlying diseases such as uncontrolled diabetes, uncontrolled blood pressure, liver and heart disease and kidney, heart and liver failure History of any of the diseases including hepatitis B or C, HIV, autoimmune diseases and immunodeficiency History of hepatocellular disease, liver failure and heart failure of liver tests more than 3 times normal History of allergy to antiviral drugs Existence of thrombocytopenia (platelets under 100000) History of recent COVID-19 vaccination lesser than 7 days being pregnant breastfeeding		Ethics committees 1 Ethics committee Name of ethics committee Ethics committee of Baqiyatallah University of Medical Sciences Street address Research center; Baqiyatallah University of Medical Sciences; Sheikh-e-bahaei street City Tehran Province Tehran Postal code 1435916471 Approval date 2021-11-20, 1400/08/29 Ethics committee reference number IR.BMSU.REC.1400.085
Age From 18 years old		Health conditions studied 1 Description of health condition studied COVID-19 ICD-10 code U07.1 ICD-10 code description COVID-19, virus identified
Gender Both		Primary outcomes 1 Description Negative pcr test or increase in CT NUMBER after intervention Timepoint Before and after the intervention Method of measurement According to the throat sample of patients with Covid-19, which is taken from patients in the same conditions in the specialized virology laboratory of Baqiyatallah Medical Center.
Phase 2-3		2 Description fever Timepoint Upon arrival and after five days
Groups that have been masked - Participant - Outcome assessor		
Sample size Target sample size: 500		
Randomization (investigator's opinion) Randomized		
Randomization description In this study, the Block Randomization method will be used using blocks of four in random order. In this method, treatment blocks are moved back and forth randomly, for example, with a block of length 4 and with two treatment groups, there are the following possibilities ABAB, ABAB, ABBA, BABA, BAAB, BBAA By putting these blocks together randomly, a balanced random list of two treatment groups is obtained		
Blinding (investigator's opinion) Double blinded		
Blinding description In this study, patients participating in the clinical trial and individuals collecting patient data will not be aware of the intervention groups. Molnupiravir and placebo will be exactly the same, and no one will know the contents until the end of the study.		
Placebo Used		

Method of measurement

with a standard thermometer

3**Description**

serum CRP inflammatory factor

Timepoint

Upon arrival and after five days

Method of measurement

By a valid laboratory

4**Description**

The level of oxygen saturation in the blood of patients

Timepoint

Upon arrival and after five days

Method of measurement

By standard pulse oximeter

5**Description**

dry cough existence

Timepoint

Upon arrival and after five days

Method of measurement

patient report and researcher observe

6**Description**

dyspnea

Timepoint

Upon arrival and after five days

Method of measurement

patient report and researcher observe

7**Description**

muscular pain

Timepoint

Upon arrival and after five days

Method of measurement

patient report and researcher observe

8**Description**

pulse rate

Timepoint

Upon arrival and after five days

Method of measurement

researcher measure

9**Description**

respiratory rate

Timepoint

Upon arrival and after five days

Method of measurement

researcher measure

10**Description**

blood pressure

Timepoint

Upon arrival and after five days

Method of measurement

researcher measure

11**Description**

serum ESR inflammatory factor

Timepoint

Upon arrival and after five days

Method of measurement

By a valid laboratory

Secondary outcomes

empty

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

Intervention group: The group receiving oral molnupiravir 200 mg daily for up to five days after the intervention, with diphenhydramine syrup / vitamin C tablet / vitamin D tablet / famotidine 40 mg tablet every 12 hours (in case of fever and pain Muscle naproxen 250 mg daily or twice daily will be added)

Category

Treatment - Drugs

2**Description**

Control group: This group will receive placebo exactly similar to molnupiravir for five days, with diphenhydramine syrup, vitamin C tablets, vitamin D tablets, famotidine 40mg tablets every 12 hours, and naproxen 250 in case of fever and muscle aches. They will receive it once or twice a day

Category

Placebo

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

Baqiyatallah hospital

Full name of responsible person

Morteza Izadi

Street address

Molasadra 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
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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
Bagheiat-allah 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
Bagheiat-allah University of Medical Sciences
Full name of responsible person
Alireza Soleymanitabar
Position
Medical student, Baqiyatallah University of Medical Sciences
Latest degree
A Level or less
Other areas of specialty/work

Person responsible for scientific inquiries

Contact

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Person responsible for updating data

Contact

Name of organization / entity
Students research committee; Baqiyatallah University of Medical Sciences
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Latest degree
A Level or less
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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

The data can be used by others without any personal information of patients

When the data will become available and for how long

after data gathering completion

To whom data/document is available

Only people with relevant permissions to use confidential information

Under which criteria data/document could be used

The data can be used for statistical analysis and re-examination

From where data/document is obtainable

Morteza Izadi; access through email:

morteza_izadi@yahoo.com

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

After calling or sending an email, the data will be sent if the applicant is valid

Comments