

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

05 Jun 2026

### Evaluation of the efficacy and safety of Favipiravir and Interferon beta compared to Lopinavir/Ritonavir and Interferon beta in patients with COVID-19

#### Protocol summary

##### Study aim

Evaluation of the efficacy and safety of Favipiravir and Interferon beta compared to Lopinavir/Ritonavir and Interferon beta in patients with COVID-19

##### Design

A phase 3 clinical trial with parallel group, open-label, 60 patients, block randomized method

##### Settings and conduct

This study will be conducted at the Shahid Mohammadi Hospital, Hormozgan University of Medical Sciences, Bandar Abbas. The study population is 60 patients with COVID-19 (30 patients in control group and 30 in study group).

##### Participants/Inclusion and exclusion criteria

Inclusion criteria: Age  $\geq 18$  years, positive polymerase chain reaction (PCR) test for COVID-19, primary clinical symptoms, hospitalized, and signing informed consent. Exclusion Criteria: Underlying diseases, including chronic hepatitis, cirrhosis, cholestatic liver diseases, cholecystitis, and peptic ulcers, acute and chronic renal failure, peptic ulcers, history of allergy to studied drugs, and pregnancy and breastfeeding.

##### Intervention groups

Group A will be patients receiving Favipiravir and Interferon beta-1a, and Group B will be patients receiving Lopinavir/Ritonavir and Interferon beta-1a.

##### Main outcome variables

Checking the viral load, fever, O<sub>2</sub> saturation, Evaluation of duration of hospitalization, C-reactive protein, Occurrence of adverse drug reactions

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20200506047323N3**

Registration date: **2020-07-22, 1399/05/01**

Registration timing: **prospective**

Last update: **2020-07-22, 1399/05/01**

Update count: **0**

##### Registration date

2020-07-22, 1399/05/01

##### Registrant information

###### Name

Mohammad Fathalipour

###### Name of organization / entity

###### Country

Iran (Islamic Republic of)

###### Phone

+98 76 3371 0406

###### Email address

m.fathalipour@hums.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2020-07-25, 1399/05/04

##### Expected recruitment end date

2020-09-25, 1399/07/04

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

##### Scientific title

Evaluation of the efficacy and safety of Favipiravir and Interferon beta compared to Lopinavir/Ritonavir and Interferon beta in patients with COVID-19

##### Public title

The effects of Favipiravir and Lopinavir/Ritonavir in treatment COVID-19

## Purpose

Treatment

## Inclusion/Exclusion criteria

### Inclusion criteria:

Age  $\geq 18$  years Positive polymerase chain reaction (PCR) test for COVID-19 Primary clinical symptoms Hospitalized Signing informed consent and willingness of study participant to accept randomization to any assigned treatment arm

### Exclusion criteria:

Underlying diseases, including chronic hepatitis, cirrhosis, cholestatic liver diseases, cholecystitis, and peptic ulcers Acute and chronic renal failure Peptic ulcers History of allergy to studied drugs Pregnancy and breastfeeding

## Age

From **18 years** old

## Gender

Both

## Phase

3

## Groups that have been masked

*No information*

## Sample size

Target sample size: **60**

## Randomization (investigator's opinion)

Randomized

## Randomization description

Block randomization will be performed (each block consists 6 patients). Allocation sequence and concealment codes will be generated using [www.sealedenvelope.com](http://www.sealedenvelope.com). The closed envelope method will be used to hide the allocation sequence.

## 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 Hormozgan University of Medical Sciences

##### Street address

Jomhuri Eslami Blvd

##### City

Bandar Abbas

##### Province

Hormozgan

## Postal code

7919915519

## Approval date

2020-07-21, 1399/04/31

## Ethics committee reference number

IR.HUMS.REC.1399.225

## Health conditions studied

### 1

#### Description of health condition studied

COVID-19 disease

#### ICD-10 code

U07.2

#### ICD-10 code description

COVID-19, virus not identified

## Primary outcomes

### 1

#### Description

Viral load

#### Timepoint

Before intervention and day 7 after the intervention

#### Method of measurement

Polymerase chain reaction (PCR) test

### 2

#### Description

Body temperature

#### Timepoint

Before intervention and daily during the study

#### Method of measurement

Thermometer

### 3

#### Description

Oxygen saturation

#### Timepoint

Before intervention and daily during the study

#### Method of measurement

Pulse oximeter

## Secondary outcomes

### 1

#### Description

Duration of hospitalization

#### Timepoint

Time period from admission to discharge

#### Method of measurement

Patient's file

### 2

#### Description

C-reactive protein

**Timepoint**

Before intervention and day 7 after the intervention

**Method of measurement**

C-RP kit

**3****Description**

Incidence of serious adverse events

**Timepoint**

Before intervention and daily during the study

**Method of measurement**

Questionnaire

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

Intervention group: Favipiravir (Zhejiang Hisun, China) plus Interferon beta-1a (CinnaGen, Iran). This group will receive 1600 mg Favipiravir twice a day for the first day and 600 mg twice a day for the following four days, plus five doses of 44 mcg Interferon beta-1a every other day. Other supportive and routine care will be the same in both groups.

**Category**

Treatment - Drugs

**2****Description**

Control group: Lopinavir/Ritonavir (Heterd company, India) plus Interferon beta-1a (CinnaGen, Iran). This group will receive 200/50 mg Lopinavir/Ritonavir twice a day for the seven days, plus five doses of 44 mcg Interferon beta-1a every other day. Other supportive and routine care will be the same in both groups.

**Category**

Treatment - Drugs

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

Shahid Mohammadi Hospital

**Full name of responsible person**

Mahdi Hasani Azad

**Street address**

Jomhuri Eslami Blvd

**City**

Bandar Abbas

**Province**

Hormozgan

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shmh@hums.ac.ir

**Web page address**

http://shmh.hums.ac.ir/

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

Bandare-abbas University of Medical Sciences

**Full name of responsible person**

Teamur Aghamolaei

**Street address**

Jomhuri Eslami Blvd

**City**

Bandar Abbas

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**Email**

mail@hums.ac.ir

**Web page address**

http://hums.ac.ir/

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

Yes

**Title of funding source**

Bandare-abbas 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**

Bandare-abbas University of Medical Sciences

**Full name of responsible person**

Mohammad Fathalipour

**Position**

Assistant professor

**Latest degree**

Ph.D.

**Other areas of specialty/work**

Medical Pharmacy

**Street address**

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

Bandare-abbas University of Medical Sciences

**Full name of responsible person**

Mohammad Fathalipour

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**Person responsible for updating data****Contact****Name of organization / entity**

Bandare-abbas University of Medical Sciences

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**Email**

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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**

Undecided - It is not yet known if there will be a plan to make this available

**Clinical Study Report**

Not applicable

**Analytic Code**

Not applicable

**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**

Information regarding the study outcomes will be shared

**When the data will become available and for how long**

Data will become available after publication of obtained results

**To whom data/document is available**

Only academic institutions

**Under which criteria data/document could be used**

The study protocol or proposal should be approved by Ethics committee of institutions. The rights of authors and sponsors should be respected.

**From where data/document is obtainable**

M.fathalipour@yahoo.com, and M.fathalipour@hums.ac.ir

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

Requests should be addressed to the Technology and Research Vice-chancellery of Hormozgan University of Medical Sciences and the project executor should be informed.

**Comments**