

Clinical Trial Protocol

Iranian Registry of Clinical Trials

28 Jun 2026

Comparison of the effect of interferon-beta 1a and Dexamethasone in hospitalized patients with Covid - 19.

Protocol summary

Study aim

Determination and comparison of the effect of beta interferon and dexamethasone in the treatment of patients with Covid-19

Design

A single-blind randomized clinical trial with a control group and parallel groups was performed on 160 patients. Block randomization method was used for randomization.

Settings and conduct

This study is being conducted at Shahid Jalil educational Hospital in Yasuj. Routine drugs with the possibility of affecting Covid 19 symptoms were used in both intervention 1 (Combination of Remdesivir, Interferon, and Dexamethasone) and intervention 2 or Control (Remdesivir and Dexamethasone) groups.

Participants/Inclusion and exclusion criteria

1. Definitive diagnosis of covid- 19 by infectious disease specialist and a positive PCR and spiral CT scan of the lung. 2. Inpatients who are at least 18 years old. 3. Hospitalized patients with pulmonary involvement in CT Scan if not intubated and not ventilated with NIV mask. 4. No history of pancytopenia. 5. Patients with a GFR of more than 30%. 6. Patients who are not pregnant. 8. Absence of liver inflammation.

Intervention groups

The intervention group1 includes patients who will receive a combination of Remdesivir, Interferon, and Dexamethasone, and the intervention group2 (Control) includes patients who will receive a combination of Remdesivir, and Dexamethasone.

Main outcome variables

The length of hospital stay Patient mortality. O2 saturation

General information

Reason for update

Acronym

مقایسه داروی بتا اینترفرون و دگزامتازون

IRCT registration information

IRCT registration number: **IRCT20150622022869N9**
Registration date: **2022-07-02, 1401/04/11**
Registration timing: **retrospective**

Last update: **2022-07-02, 1401/04/11**

Update count: **0**

Registration date

2022-07-02, 1401/04/11

Registrant information

Name

Moslem Sedaghattalab

Name of organization / entity

Yasuj University of Medical Sciences

Country

Iran (Islamic Republic of)

Phone

+98 74 3322 0163

Email address

m.sedaghattalab@yums.ac.ir

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2021-09-23, 1400/07/01

Expected recruitment end date

2022-06-22, 1401/04/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Comparison of the effect of interferon-beta 1a and Dexamethasone in hospitalized patients with Covid - 19.

Public title

Comparison of Interferon-beta-1a and Dexamethasone on hospitalized patients with Covid-19.

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria:

Definitive diagnosis of covid-19 by infectious disease specialist with a positive PCR and spiral ct scan of the lung. Hospitalized patients who are at least 18 years old.

Exclusion criteria:

Pregnant women Patients with GFR less than 30%. Patient with a history of Pancytopenia. Increased ALT more than 10 times.

Age

From **18 years** old

Gender

Both

Phase

3

Groups that have been masked

- Outcome assessor

Sample size

Target sample size: **176**

Randomization (investigator's opinion)

Randomized

Randomization description

Block randomization method will be used. A total of 176 patients with Covid-19 who meet the inclusion criteria will be selected and randomly assigned to the two groups of intervention 1 and intervention 2. In block randomization, according to the total number of 176 patients and the existence of two groups, blocks with sizes of 2, 4, and 6 will be used. For example, if the size of block is 2, we will have two modes AB and BA (A and B are intervention groups 1 and intervention 2). In order to create sequences and concealment, online software with the following address will be used, and a special randomization code will be generated. Sealed Envelope Ltd. 2021. Create a blocked randomisation list. [Online] Available from: <https://www.sealedenvelope.com/simple-randomiser/v1/lists> [Accessed 5 Jun 2022]

Blinding (investigator's opinion)

Single blinded

Blinding description

In this study, the person evaluating the outcome variables is unaware of the study groups.

Placebo

Not used

Assignment

Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

کمیته اخلاق دانشگاه علوم پزشکی یاسوج

Street address

Shahid Motahari Boulevard

City

Yasuj

Province

Kohgilouyeh-va-Boyerahmad

Postal code

7591874934

Approval date

2021-09-21, 1400/06/30

Ethics committee reference number

IR.YUMS.REC.1400.119

Health conditions studied

1

Description of health condition studied

COVID - 19

ICD-10 code

U07.1

ICD-10 code description

COVID-19, virus identified

Primary outcomes

1

Description

Arterial oxygen saturation

Timepoint

At the time of admission and discharge of the patient

Method of measurement

pulse oximeter device

Secondary outcomes

1

Description

Mortality rate

Timepoint

At the end of the intervention and completion of sampling

Method of measurement

By comparing the number of deaths in each group

2

Description

Duration of hospitalization

Timepoint

On the day of discharge from the hospital

Method of measurement

Number of hospitalization days

Intervention groups

1

Description

Intervention group: (combination of three drugs Remdesivir, Interferon, and Dexamethasone) Remdesivir drugs with a dose of 200 mg in the initial dose and then 100 mg daily for 5 days, Interferon-beta 1 alpha is given as a subcutaneous injection every other day for 4 doses (each vial contains 30 micro-grams of interferon is equivalent to 6 million international units) and dexamethasone at a dose of 8 mg daily intravenously.

Category

Treatment - Drugs

2

Description

Intervention group 2: Combination of Remdesivir and Dexamethasone) Remdesivir with a dose of 200 mg in the initial dose and then 100 mg daily for 5 days, and Dexamethasone at a dose of 8 mg daily intravenously.

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Shahid Jalil Hospital in Yasuj

Full name of responsible person

Moslem Sedaghattalab

Street address

Shahid Qharani Boulevard

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Kohgiluyeh-va-Boyrahmad

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M.sedaghattalab@yums.ac.ir

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Yasouj University of Medical Sciences

Full name of responsible person

Moslem Sedaghattalab

Street address

بلوار شهید مطهری

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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

Yasuj 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

Yasouj University of Medical Sciences

Full name of responsible person

Moslem Sedaghattalab

Position

Assistant Professor

Latest degree

Specialist

Other areas of specialty/work

Internal Medicine

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Person responsible for scientific inquiries

Contact

Name of organization / entity

Yasouj University of Medical Sciences

Full name of responsible person

M.sedaghattalab@yums.ac.ir

Position

Assistant Professor

Latest degree

Specialist

Other areas of specialty/work

Internal Medicine

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

Yasouj University of Medical Sciences

Full name of responsible person

Moslem Sedaghattalab

Position

Assistant Professor

Latest degree

Specialist

Other areas of specialty/work

Internal Medicine

Street address**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 more information.

Study Protocol

No - There is not a plan to make this available

Statistical Analysis Plan

No - There is not 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

No - There is not a plan to make this available

Analytic Code

No - There is not a plan to make this available

Data Dictionary

No - There is not a plan to make this available