

Clinical Trial Protocol

Iranian Registry of Clinical Trials

13 Jun 2026

Comparative study of the therapeutic effect of citrizine and fexofenadine in the control of medication-induced skin lesions in hospital intensive care unit

Protocol summary

Study aim

Comparative study of the efficacy of citrizine and fexofenadine in controlling drug-induced skin lesions in hospital intensive care unit to control skin lesions and improve patient quality of life and reduce mortality in hospital intensive care unit

Design

Clinical trial on sixty patients in two groups, community-based and pragmatic therapy, with parallel groups, single blind, randomized to two groups

Settings and conduct

Patients admitted to the intensive care unit will receive either oral or gavage Fexofenadine tablets 120 mg every 12 hours for a maximum of ten days and oral or gavage cetirizine tablets 10 mg every 12 hours. Then questionnaire information before and after It will be completed after receiving the drug. unilateral blinding is done so that the participant is blind and does not know the grouping and the number of participants in the study.

Participants/Inclusion and exclusion criteria

Obtain informed and patient consent in the ICU for at least 48 hours. No history of skin diseases.

Intervention groups

Fexofenadine 120 mg of one tablet every 12 hour for ten days in one group and cetirizine 10 mg of one tablet every 12 hour for two week in another group

Main outcome variables

Evaluation of drug-induced skin lesions; severity of drug-induced skin lesions and extent of drug-induced skin lesions

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20080901001165N47**

Registration date: **2020-04-16, 1399/01/28**

Registration timing: **retrospective**

Last update: **2020-04-16, 1399/01/28**

Update count: **0**

Registration date

2020-04-16, 1399/01/28

Registrant information

Name

Yunes Panahi

Name of organization / entity

Baqiyatallah University of Medical Sciences

Country

Iran (Islamic Republic of)

Phone

+98 21 8821 1524

Email address

yunespanahi@bmsu.ac.ir

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2018-12-01, 1397/09/10

Expected recruitment end date

2019-11-06, 1398/08/15

Actual recruitment start date

2018-12-11, 1397/09/20

Actual recruitment end date

2020-02-09, 1398/11/20

Trial completion date

2020-02-19, 1398/11/30

Scientific title

Comparative study of the therapeutic effect of citrizine and fexofenadine in the control of medication-induced skin lesions in hospital intensive care unit

Public title

A comparative study of the therapeutic effect of cetirizine and fexofenadine on the control of skin lesions caused by drugs in the intensive care unit(ICU).

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria:

Patients stay in ICU for at least 48 hours No history of skin diseases Be on medication, so that people in both groups receive their own treatment appropriate to their illness.

Exclusion criteria:

The patient is in the ICU for less than 48 hours. Find non-pharmacological skin diseases

Age

From **18 years** old

Gender

Both

Phase

N/A

Groups that have been masked

- Participant

Sample size

Target sample size: **60**

Actual sample size reached: **80**

Randomization (investigator's opinion)

Randomized

Randomization description

Randomization is conducted using random number table and considering a 1:1 ratio.

Blinding (investigator's opinion)

Single blinded

Blinding description

The patient groups are divided into two groups and except the researcher, other people involved in the study do not know of the intervention or control group.

Placebo

Not used

Assignment

Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethics Committee of Islamic Azad University /
Pharmaceutical Sciences Branch

Street address

Yasaman alley, Yakhchal steet, Shariati street

City

Tehran

Province

Tehran

Postal code

1941933111

Approval date

2018-08-28, 1397/06/06

Ethics committee reference number

IR.IAU.PS.REC.1397.348

Health conditions studied

1

Description of health condition studied

Drug induced skin eruption

ICD-10 code

R21

ICD-10 code description

Rash and other nonspecific skin eruption

Primary outcomes

1

Description

Type of drug-induced skin lesion

Timepoint

Before intervention, 10 days after intervention

Method of measurement

Questionnaire

2

Description

The severity of the skin lesions

Timepoint

Before intervention, 10 days after intervention

Method of measurement

questionnaire

3

Description

The extent of skin lesions

Timepoint

Before intervention, 10 days after intervention

Method of measurement

questionnaire

Secondary outcomes

1

Description

Side effect of Faxofenadine

Timepoint

Before intervention, after ten days

Method of measurement

Questionnaire

2

Description

Side effect of cetirizine

Timepoint

Before intervention, after ten days

Method of measurement

Questionnaire

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

Intervention group: Fexofenadine tablets 120 mg every 12 hours for 10 days

Category

Treatment - Drugs

2**Description**

Intervention group: Cetirizine 10 mg tablet every 12 hours for 10 days

Category

Treatment - Drugs

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

Baqiyatallah Specialty Hospital

Full name of responsible person

Yunes panahi

Street address

Wank-Sheikh Baha'i South street -Baqiyatallah
Faculty of Pharmacy

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

Islamic Azad University

Full name of responsible person

Farshad Hashemian

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Islamic Azad University of Pharmaceutical Sciences,
Yakhchal St., Shariati St., Tehran, Iran

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Hashemian.f@iaups.ac.ir

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

Yes

Title of funding source

Islamic Azad University

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

Yunes panahi

Position

pharmacotherapy professor

Latest degree

Specialist

Other areas of specialty/work

Medical Pharmacy

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

Bagheiat-allah University of Medical Sciences

Full name of responsible person

Yunes panahi

Position

pharmacotherapy professor

Latest degree

Specialist

Other areas of specialty/work

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

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Full name of responsible person

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Position

pharmacy student

Latest degree

A Level or less

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

No more information

Study Protocol

No - There is not a plan to make this available

Statistical Analysis Plan

Not applicable

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

Not applicable

Data Dictionary

Not applicable

Title and more details about the data/document

The clinical report will be published in paper form

When the data will become available and for how long

Start access period after printing results

To whom data/document is available

Researchers in the field of health

Under which criteria data/document could be used

If the documentation is protected

From where data/document is obtainable

yunes panahi

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

Send project email

Comments