

Clinical Trial Protocol

Iranian Registry of Clinical Trials

05 Jul 2026

Evaluation the clinical effects of short -course herbal drug consist of Honey and zufa on Asthma symptoms and Exacerbation due to viral respiratory infection in Children with mild intermittent Asthma

Protocol summary

Summary

This study as a three blind clinical trial in two groups of intervention and control was completed. 50 children of 7 to 12 years old with the diagnose of mild intermittent asthma by the specialists of Immunology and Allergy were enrolled. The children were randomly assigned to two experimental and control groups. Patients at the beginning of a common cold by expert were examined and history, physical examination and peak flow expiratory using a peak flow meter were recorded, then herbal medicine (combined with honey and extracts of hyssop, mallow, thyme narrow, violets and Khorasani dates, dried figs, licorice, mallow, veneris) was administrated to the intervention group and placebo (a substance like herbs with the same shape, odor, design and color) was administered to the control group. In the intervention group patients according to age received a certain amount of medicinal plants (5 ml every 8 hours). The treatment was continued for 5 days. Again on the third day and the fourteenth day by phone and in seventh day in personal were examined and the information (history, physical examination and peak flow) were noted in the checklist.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT201608314976N5**
Registration date: **2017-07-22, 1396/04/31**
Registration timing: **retrospective**

Last update:

Update count: **0**

Registration date

2017-07-22, 1396/04/31

Registrant information

Name

Hamid Ahanchian

Name of organization / entity

Mashhad University of Medical Sciences

Country

Iran (Islamic Republic of)

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+98 51 3801 2469

Email address

ahanchianh@mums.ac.ir

Recruitment status

Recruitment complete

Funding source

vice chancellor for research, Mashhad University of medical sciences.

Expected recruitment start date

2015-09-23, 1394/07/01

Expected recruitment end date

2016-12-20, 1395/09/30

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Evaluation the clinical effects of short -course herbal drug consist of Honey and zufa on Asthma symptoms and Exacerbation due to viral respiratory infection in Children with mild intermittent Asthma

Public title

effect of herbal drug on respiratoty infection in asthmatic children

Purpose

Treatment

Inclusion/Exclusion criteria

inclusion criteria: age of 7 to 12 years, mild to moderate asthma, acute viral upper respiratory tract infection, absence of congenital anomalies, absence of any other disease. exclusion criteria: bacterial infection, drug reaction, inadequate compliance of patient or family in follow up or drug consumption.

Age

From **7 years** old to **12 years** old

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **50**

Randomization (investigator's opinion)

Randomized

Randomization description

Blinding (investigator's opinion)

Triple blinded

Blinding description

Placebo

Used

Assignment

Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethic committe of Mashhad university of Medical Sciences

Street address

Ghoreyshi Building, Daneshgah street

City

mashhad

Postal code

Approval date

2016-01-06, 1394/10/16

Ethics committee reference number

IR.MUMS.FM.REC.1394.374

Health conditions studied

1

Description of health condition studied

asthma

ICD-10 code

J44

ICD-10 code description

chronic obstructive asthma

Primary outcomes

1

Description

cough

Timepoint

0 and 3rd 7th and 14teen days

Method of measurement

numbers

2

Description

awaking during sleep

Timepoint

day 0,3,7,14

Method of measurement

number

Secondary outcomes

1

Description

PEFR (Peak Expirometry Flow Rate) changes

Timepoint

0 3rd days

Method of measurement

Peak Expirometry Flow Rate device

2

Description

needing to adjuant treatment (montelukast)

Timepoint

day3,7,14

Method of measurement

yes or no

Intervention groups

1

Description

interbention group: Compounds containing honey and hyssop herb extracts, hibiscus, thyme narrow, and jujube Khorasani violets, dried figs, licorice, mallow, veneris witch was produced by pharmacist professor in Emam-Reza pharmacy and with dose of 5 milliliter every 8 hour were prescribed.

Category

Treatment - Drugs

2

Description

control group: placebo (a substance like herbs with the same shape,Odor, design and color) witch was produced by pharmacist professor in Emam-Reza pharmacy and with dose of 5 milliliter every 8 hour were prescribed.

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Ghaem Hospital

Full name of responsible person

Dr.hamid ahanchian

Street address

City

Mashhad

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

mashhad university of medical science

Full name of responsible person

DR mohsen tafaghodi

Street address

mashhad daneshgah street, ghoreishi beelding, vice
chancellor for research of mashhad university of
medical science

City

Mashhad

Grant name

-

Grant code / Reference number

-

Is the source of funding the same sponsor organization/entity?

Yes

Title of funding source

mashhad university of medical science

Proportion provided by this source

100

Public or private sector

empty

Domestic or foreign origin

empty

Category of foreign source of funding

empty

Country of origin

Type of organization providing the funding

empty

Person responsible for general inquiries

Contact

Name of organization / entity

Mashhad University of Medical Sciences

Full name of responsible person

Hamid Ahanchian

Position

Associate professor

Other areas of specialty/work

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Sharing plan

Deidentified Individual Participant Data Set (IPD)

empty
Study Protocol
empty
Statistical Analysis Plan
empty
Informed Consent Form
empty

Clinical Study Report
empty
Analytic Code
empty
Data Dictionary
empty