

Clinical Trial Protocol

Iranian Registry of Clinical Trials

03 Jul 2026

Comparison of ferrous sulfate with ferrous sulfate and traditional medicine product (currant and pomegranate) effect on regulation of ferritin in iron deficient women with normal hemoglobin

Protocol summary

2015-12-02, 1394/09/11

Summary

Target: Comparison of ferrous sulfate with ferrous sulfate and traditional medicine product (currant and pomegranate) effect on regulation of ferritin in iron deficient women with normal hemoglobin. Design study: accidental and no blindness patient and researcher on the project. The intervention group will be compared with the control group. Inclusion criteria: Hemoglobin above 12, ferritin below 50 Exclusion criteria: cardiovascular disease; thyroid or renal or hepatic disease; use supplement or Iron; hormone therapy; AUB; GI bleeding; abnormal CRP; smoking; menopause; pregnancy; and the factors related to abnormal red blood cells. Intervention group will receive traditional product (currant and pomegranate) once a day at fast, plus ferrous sulfate up to three tablets per day. Control group will receive ferrous sulfate, up to three tablets per day. Follow up: every 3 month up to 9 month with control blood Tests, and monthly by phone. Major outcome: the main outcome of the project is modified ferritin. Measurement of CBC, CRP every three months, and on three occasions (up to 9 months) of patients is done. At each period, people treated themselves and Speed-called ferritin levels in the two groups will be compared with each other.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2015083123835N1**
Registration date: **2015-12-02, 1394/09/11**
Registration timing: **registered_while_recruiting**

Last update:

Update count: **0**

Registration date

Registrant information

Name

Somaye Fatali

Name of organization / entity

Kerman University of Medical Sciences

Country

Iran (Islamic Republic of)

Phone

+98 21 2249 5995

Email address

s-fatali@razi.tums.ac.ir

Recruitment status

Recruitment complete

Funding source

College of Traditional Medicine, Sajadie Ave, Modiriat Ave, Kerman

Expected recruitment start date

2015-09-11, 1394/06/20

Expected recruitment end date

2016-06-19, 1395/03/30

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Comparison of ferrous sulfate with ferrous sulfate and traditional medicine product (currant and pomegranate) effect on regulation of ferritin in iron deficient women with normal hemoglobin

Public title

Helping to treatment of iron deficiency with traditional medicine product

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria : women with HB > 12 and Ferritin < 50

Exclusion criteria: cardiovascular disease; thyroid or renal or hepatic disease; use supplement or Iron; hormone therapy; AUB; GI bleeding; abnormal CRP; smoking; menopause; pregnancy; and the factors related to abnormal red blood cells.

Age

From **18 years** old to **45 years** old

Gender

Female

Phase

2-3

Groups that have been masked

No information

Sample size

Target sample size: **74**

Randomization (investigator's opinion)

Randomized

Randomization description

Blinding (investigator's opinion)

Not blinded

Blinding description

Placebo

Not used

Assignment

Parallel

Other design features

Secondary Ids

1

Registry name

Secondary trial Id

Registration date

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethics Committee of Kerman University of Medical Sciences

Street address

Central Organization of Kerman University of Medical Sciences, Jomhuri islami Blv., Kerman

City

Kerman

Postal code

۷۶۱۵۶۸۷۱۳۴

Approval date

2015-09-09, 1394/06/18

Ethics committee reference number

ir.kmu.rec.1394.231

Health conditions studied

1

Description of health condition studied

Iron deficiency anemia

ICD-10 code

D50.9

ICD-10 code description

Iron deficiency anaemia, unspecified

Primary outcomes

1

Description

Ferritin

Timepoint

3 month

Method of measurement

Blood tests

2

Description

Hemoglobin

Timepoint

3 month

Method of measurement

Blood tests

Secondary outcomes

1

Description

Constipation

Timepoint

3 month

Method of measurement

The number of bowel movements per day

Intervention groups

1

Description

Intervention group: traditional product(currant and pomegranate) in fasting, plus ferrous sulfate up to three tablets per day

Category

Treatment - Drugs

2

Description

Control group: ferrous sulfate , up to three tablets per day

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Imam Khomainsi Hospital, Hematological Clinic

Full name of responsible person

Somaye Fatali

Street address

Gharib Ave, Keshavars Blv, Tehran

City

Tehran

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Vice chancellor for research, Kerman University of Medical Sciences

Full name of responsible person

Hale Tajoddini

Street address

College of Traditional Medicine, Sajadieh Ave, Modiriat Ave, Kerman

City

Kerman

Grant name

Grant code / Reference number

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

Yes

Title of funding source

Vice chancellor for research, Kerman University of Medical Sciences

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

Kerman University of Medical Sciences

Full name of responsible person

Somaye Fatali

Position

Ph.D Student of Iranian Traditional Medicine

Other areas of specialty/work

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Person responsible for scientific inquiries

Contact

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Full name of responsible person

Mitra Mehrabani

Position

Pharmacognosist

Other areas of specialty/work

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Position

Ph.D student of Iranian Traditional Medicine

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Email

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Web page address

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