

Clinical Trial Protocol

Iranian Registry of Clinical Trials

26 May 2026

The study of effect of selenium on acute phase reactants and thyroid function test in hemodialysis patients

Protocol summary

Summary

Objectives: the aim of this study is understanding the effect of selenium on acute phase reactants and thyroid function test in hemodialysis patients. Design: randomised clinical trial. Setting and conduct: after determining the serum level of selenium in hemodialysis patients who referred to Emam Reza hospital, 100 patients who have selenium deficiency divided into 2 groups by age, sex and body mass index. One group treated by selenium capsule 200mg daily for 3 months and the other group treated by placebo capsule. Before and after intervention we measured ESR, CRP, FERRITIN, TSH, T4 and T3RU. Participants including major eligibility criteria: Inclusion criteria: the patient's willingness to cooperate; at least 6 months passing after dialysis starting . Exclusion criteria: infectious diseases during last 3 months; steroidal and non-steroidal anti-inflammatory drugs; vitamins E,C and zinc usage within two months before the study. Interventions: selenium capsule 200 mg daily for 3 months. Placebo capsule daily for 3 months. Main outcome measure: ESR, CRP, FERRITIN, TSH, T4, T3RU

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2014041517284N1**

Registration date: **2014-05-20, 1393/02/30**

Registration timing: **retrospective**

Last update:

Update count: **0**

Registration date

2014-05-20, 1393/02/30

Registrant information

Name

Kanan Nikseresht

Name of organization / entity

Kermanshah University of medical sciences

Country

Iran (Islamic Republic of)

Phone

+98 83 1427 6300

Email address

kanan-nik@kums.ac.ir

Recruitment status

Recruitment complete

Funding source

Vice Chancellor for Research Affairs, Kermanshah University Of Medical Sciences

Expected recruitment start date

2013-03-20, 1391/12/30

Expected recruitment end date

2014-03-20, 1392/12/29

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The study of effect of selenium on acute phase reactants and thyroid function test in hemodialysis patients

Public title

The study of effect of selenium on acute phase reactants and thyroid function test in hemodialysis patients

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria: the patient's willingness to cooperate; at least 6 months passing after dialysis starting .
Exclusion criteria: infectious diseases during last 3 months; steroidal and non-steroidal anti-inflammatory drugs; vitamins E, C and zinc usage within two months

before the study.

Age

No age limit

Gender

Both

Phase

2-3

Groups that have been masked

No information

Sample size

Target sample size: **100**

Randomization (investigator's opinion)

N/A

Randomization description**Blinding (investigator's opinion)**

Double blinded

Blinding description**Placebo**

Used

Assignment

Parallel

Other design features**Secondary Ids****1****Registry name**

Not registered

Secondary trial Id

Not registered yet

Registration date

empty

Ethics committees**1****Ethics committee****Name of ethics committee**

Ethics committee of Kermanshah University of
Medical Sciences

Street address

Building No.2, Shahid Beheshti Blvd, Vice Chancellor
for Research Affairs, Kermanshah University Of
Medical Sciences

City

Kermanshah

Postal code**Approval date**

2013-09-23, 1392/07/01

Ethics committee reference number

۴۹۴۰۱ شماره نامه/۷/۴۲۰/پ

Health conditions studied**1****Description of health condition studied**

Hemodialysis patient

ICD-10 code

N18.0

ICD-10 code description

End-Stage renal disease

Primary outcomes**1****Description**

Selenium

Timepoint

90 days

Method of measurement

By cell counter

Secondary outcomes**1****Description**

Quantity CRP

Timepoint

3 months later

Method of measurement

Cell counter

2**Description**

T3RU

Timepoint

3months later

Method of measurement

Cell counter

3**Description**

T4

Timepoint

3months later

Method of measurement

Cell counter

4**Description**

TSH

Timepoint

3months later

Method of measurement

Cell counter

5**Description**

ESR

Timepoint

3months later

Method of measurement

Cell counter

6

Description

Ferritin

Timepoint

3months later

Method of measurement

Cell counter

Intervention groups

1

Description

Daily prescription of selenium supplement for 3 months

Category

Treatment - Drugs

2

Description

Daily prescription of placebo capsule containing starch for 3 months

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Emam Reza hospital, Kermanshah

Full name of responsible person

Dr. Kanan Nikseresht

Street address

Internal ward, Emam Reza hospital, Zakaria razi Blvd.

City

Kermanshah

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Vice Chancellor for Research Affairs, Kermanshah University Of Medical Sciences

Full name of responsible person

Koroush Hamzehee

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Building No.2, Shahid Beheshti, Vice Chancellor for Research Affairs, Kermanshah University Of Medical Sciences

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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 Affairs, Kermanshah 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

Kermanshah University of Medical Sciences

Full name of responsible person

Dr. Kanan Nikseresht

Position

Internal medicine resident

Other areas of specialty/work

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Person responsible for updating data

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Name of organization / entity

Kermanshah University of Medical Sciences

Full name of responsible person

Kanan Nikseresht

Position

Internal medicine resident

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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