

Clinical Trial Protocol

Iranian Registry of Clinical Trials

09 Jun 2026

Effect of vitamin D supplementation on metabolic status, thyroid hormones, serum irisin and obesity indices in women with subclinical hypothyroidism

Protocol summary

Study aim

Effect of vitamin D supplementation on metabolic status, thyroid hormones, serum irisin and obesity indices in women with subclinical hypothyroidism

Design

A randomized, double-blind, placebo-controlled trial

Settings and conduct

44 patients with subclinical hypothyroidism will be randomly divided into two groups: the group receiving vitamin D (one per 50,000 units per week) and the group receiving placebo (one per 1 containing sunflower oil similar to vitamin D). The patients will be selected from the endocrinology clinic of Imam Reza Hospital in Tabriz.

Participants/Inclusion and exclusion criteria

A total of 44 patients with hypothyroidism will be selected based on the inclusion criteria and informed written consent will be obtained from all participants before enrollment. Inclusion criteria: willingness to participate in the study, women aged 20 to 65 years with hypothyroidism, BMI 25-37 Kg / m², laboratory TSH between 5-10 micro units per ml and normal T3 and T4, test Anti-TPO positive; Exclusion criteria: unwillingness to participate in the study; Serum vitamin D level higher than 100 micrograms per liter; Smoking and alcohol; Pregnancy and lactation; Diabetes and infectious heart disease; Renal failure; Cushing's syndrome; Liver disease; Use of drugs that affect the level of thyroid hormones including lithium, amiodarone, steroids, hypolipidemic drugs, antidepressants, oral contraceptives; Regular consumption of any dietary supplement during the last 2 months and during the study; Congenital hypothyroidism; Hypothyroidism due to radiation therapy to the head and neck

Intervention groups

1. Vitamin D supplement group (50000 units weekly)
2. Placebo group

Main outcome variables

glycemic indices; serum level of lipid profile; thyroid hormones; serum Irisin; anthropometric measurements; vitamin D serum

General information

Reason for update

Acronym

VITDHY POT

IRCT registration information

IRCT registration number: **IRCT20100408003664N25**

Registration date: **2021-10-23, 1400/08/01**

Registration timing: **prospective**

Last update: **2021-10-23, 1400/08/01**

Update count: **0**

Registration date

2021-10-23, 1400/08/01

Registrant information

Name

Maryam Rafrat

Name of organization / entity

Tabriz University Of Medical Sciences

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2021-11-06, 1400/08/15

Expected recruitment end date

2022-03-06, 1400/12/15

Actual recruitment start date
empty

Actual recruitment end date
empty

Trial completion date
empty

Scientific title
Effect of vitamin D supplementation on metabolic status, thyroid hormones, serum irisin and obesity indices in women with subclinical hypothyroidism

Public title
The effects of vitamin D supplementation in women with hypothyroidism

Purpose
Treatment

Inclusion/Exclusion criteria
Inclusion criteria:
Willingness to participate in research Women between the ages of 20 and 65 with subclinical hypothyroidism BMI 25-37 Kg / m² TSH 5-10 micro units per milliliter and freeT3 and free T4 are normal
Exclusion criteria:
Serum Vitamin D Levels higher than 100 µg / L Smoking and alcohol Pregnancy and lactation Diabetes and infectious heart disease Kidney failure Cushing's syndrome Liver diseases Regular consumption of any dietary supplement during the last 2 months and during the study

Age
From **20 years** old to **65 years** old

Gender
Female

Phase
3

Groups that have been masked

- Participant
- Investigator

Sample size
Target sample size: **44**

Randomization (investigator's opinion)
Randomized

Randomization description
The eligible participants will be randomly allocated to intervention and placebo groups using a software-generated random permuted blocks. The generated random sequence will be kept in a protected location and administered by an independent third party who is blind to the trial throughout the study. In order to hide, a random sequence will not be provided to the executor and will be performed sequentially for each block.

Blinding (investigator's opinion)
Double blinded

Blinding description
The placebo and supplement will be packed in the same number in similar packages. The method of blindness will be that the supplements and placebo will be delivered to the participants by someone other than the researcher, and the researcher will remain unaware until the end of

the study.

Placebo
Used

Assignment
Parallel

Other design features

Secondary Ids
empty

Ethics committees

1

Ethics committee
Name of ethics committee
Ethics Committee of Tabriz University of Medical Science
Street address
Tabriz University of Medical Science, Attar Neishabouri Avenue, Golgasht street, Tabriz
City
Tabriz
Province
East Azarbaijan
Postal code
5138663134

Approval date
2021-07-10, 1400/04/19

Ethics committee reference number
IR.TBZMED.REC.1400.327

Health conditions studied

1

Description of health condition studied
subclinical hypothyroidism

ICD-10 code
E03

ICD-10 code description
Other hypothyroidism

Primary outcomes

1

Description
Serum Vitamin D

Timepoint
Baseline and 12 weeks after the intervention

Method of measurement
ELISA

2

Description
Glycemic indexes

Timepoint
Baseline and after 12 weeks after the intervention

Method of measurement

Glycemic indexes (from indicators of metabolic status) including fasting blood sugar by enzymatic method, insulin by ELISA and insulin resistance by formula, will be measured or calculated.

3

Description

lipid profile

Timepoint

Baseline and 12 weeks after the intervention

Method of measurement

Lipid profile (from indicators of metabolic status) including total cholesterol, triglycerides and high-density lipoprotein will be measured enzymatically, and low-density lipoprotein will be calculated using the friedwald formula.

4

Description

Thyroid hormones

Timepoint

Baseline and 12 weeks after the intervention

Method of measurement

Thyroid hormones including thyroid stimulating hormone, triiodothyronine and thyroxine will be measured by ELISA.

5

Description

Irisin

Timepoint

Baseline and 12 weeks after the intervention

Method of measurement

ELISA

Secondary outcomes

1

Description

Obesity indices

Timepoint

Baseline and 12 weeks after the intervention

Method of measurement

Obesity indices including weight with scales and waist circumference and hip circumference will be measured with a tape measure. Body mass index will be calculated with the formula.

2

Description

Body composition

Timepoint

Baseline and 12 weeks after the intervention

Method of measurement

Body composition will be measured by Bio impedance (BIA) method using Tanita body composition analyzer.

3

Description

Dietary intake

Timepoint

Baseline and 12 weeks after the intervention

Method of measurement

The food record questionnaire will be used to assess food intake including energy, carbohydrate, protein and fat.

4

Description

Physical activity

Timepoint

Baseline and 12 weeks after the intervention

Method of measurement

The International Physical Activity Questionnaire will be used to measure physical activity.

Intervention groups

1

Description

Intervention group: 50000 units weekly vitamin D of Zahravi company for 3 month

Category

Treatment - Drugs

2

Description

Control group: Intake a Placebo perl containing sunflower oil, which is similar to vitamin D supplement in terms of dosage, color and size, once a week for 12 weeks

Category

Placebo

Recruitment centers

1

Recruitment center

Name of recruitment center

Endocrinology clinic of Imam Reza Hospital , Tabriz

Full name of responsible person

Sara Safari

Street address

Imam Reza Hospital, Golghast Street, Tabriz

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

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Full name of responsible person

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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

Tabriz 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**

Tabriz University of Medical Sciences

Full name of responsible person

Sara Safari

Position

MSc Student of Nutrition Sciences

Latest degree

Bachelor

Other areas of specialty/work

Nutrition

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Latest degree

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Position

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Latest degree

Bachelor

Other areas of specialty/work

Nutrition

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Sharing plan**Deidentified Individual Participant Data Set (IPD)**

Undecided - It is not yet known if there will be a plan to make this available

Study Protocol

Undecided - It is not yet known if there will be a plan to make this available

Statistical Analysis Plan

Undecided - It is not yet known if there will be 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

Undecided - It is not yet known if there will be a plan to make this available

Analytic Code

Undecided - It is not yet known if there will be a plan to make this available

Data Dictionary

Undecided - It is not yet known if there will be a plan to make this available