

Clinical Trial Protocol

Iranian Registry of Clinical Trials

08 Sep 2026

Assessment of the effect of Levothyroxine on the infant development status in pregnant women with subclinical hypothyroidism

Protocol summary

Summary

This study will be conducted in two parts: In the first stage, 1600 pregnant women were evaluated and a questionnaire including demographic characteristics, medical and midwifery history were completed. Also, their thyroid is examined. Blood samples were taken from sera for TSH, T4, T-uptake and TPOAb tests. According to the results of the laboratory test, TPOAb negative women (<50 IU/mL) with TSH 2.5–10 µIU/mL in The first 20 weeks of pregnancy is identified as sub-clinical hypothyroidism with negative thyroid peroxidase antibodies (TPOAb-) and entering the second stage of the study. In the second stage, women with subclinical hypothyroidism TPOAb- are randomly divided into two groups of intervention and control. The first group is the intervention group that is treated with levothyroxine and the second group does not receive any intervention. Pregnant women are followed to delivery. After delivery, the developmental status of children is evaluated using Ages and Stages Questionnaire (ASQ) completed by parents for each of the five areas of communication, gross motor, Fine motions, problem solving and personal-social skills. The Children developmental status in the two interventions and control groups is examined and compared.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2017090314849N5**
Registration date: **2017-09-11, 1396/06/20**
Registration timing: **prospective**

Last update:

Update count: **0**

Registration date

2017-09-11, 1396/06/20

Registrant information

Name

Sima Nazarpour

Name of organization / entity

Shahid Beheshti University of Medical Sciences

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Shahid Beheshti University of Medical Sciences

Expected recruitment start date

2017-09-23, 1396/07/01

Expected recruitment end date

2019-03-20, 1397/12/29

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Assessment of the effect of Levothyroxine on the infant development status in pregnant women with subclinical hypothyroidism

Public title

The effect of Levothyroxine on the infant development status in pregnant women with subclinical hypothyroidism

Purpose

Prevention

Inclusion/Exclusion criteria

Inclusion criteria: pregnant women during the first 20 weeks of pregnancy. Exclusion criteria: multiple

pregnancy.

Age
From **15 years** old to **46 years** old

Gender
Female

Phase
N/A

Groups that have been masked
No information

Sample size
Target sample size: **1600**

Randomization (investigator's opinion)
Randomized

Randomization description

Blinding (investigator's opinion)
Single blinded

Blinding description

Placebo
Not used

Assignment
Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee
Ethics Committee Research Institute for Endocrine Sciences and Metabolism of Shahid Beheshti Univers

Street address
No.24 Arabi street- Yemen street - Velenjak- Tehran- Iran

City
Tehran

Postal code
1985717413

Approval date
2017-03-01, 1395/12/11

Ethics committee reference number
IR.SBMU.ENDOCRINE.REC.1395.300

Health conditions studied

1

Description of health condition studied
Subclinical hypothyroidism

ICD-10 code
E02

ICD-10 code description
Subclinical iodine-deficiency hypothyroidism

Primary outcomes

1

Description
Gross motor developmental status

Timepoint
Since 4 months after birth

Method of measurement
Ages and Stages Questionnaire (ASQ)

2

Description
problem solving developmental status

Timepoint
Since 4 months after birth

Method of measurement
Ages and Stages Questionnaire (ASQ)

3

Description
Fine motor developmental status

Timepoint
Since 4 months after birth

Method of measurement
Ages and Stages Questionnaire (ASQ)

4

Description
personal - social developmental status

Timepoint
Since 4 months after birth

Method of measurement
Ages and Stages Questionnaire (ASQ)

5

Description
communication developmental status

Timepoint
Since 4 months after birth

Method of measurement
Ages and Stages Questionnaire (ASQ)

Secondary outcomes

empty

Intervention groups

1

Description
Intervention group will treat with Levoxine (Levothyroxine sodium). These patients will receive 1 mg per kg of body weight daily levothyroxine. Dosages will maintain throughout pregnancy until delivery. In this study are used from Levothyroxine sodium 0.1 mg manufactured by Iran Hormone.

Category
Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Centers under coverage of Shahid Beheshti University of Medical Sciences (Alghadir, Kahnak, Fardis,

Full name of responsible person

responsible midwife of center

Street address

City

Tehran and Varamin

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Vice Chancellor for research of Shahid Beheshti University of Medical Sciences

Full name of responsible person

Grant for research projects section

Street address

7th Floor, Bldg No.2 SBUMS, Arabi Ave, Daneshjoo Blvd, Velenjak, Tehran, Iran. Post Code: 19839-63113

City

Tehran

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 of Shahid Beheshti 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

Research Institute for Endocrine Sciences

Full name of responsible person

Sima Nazarpour

Position

PhD / Faculty member, Researcher

Other areas of specialty/work

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Person responsible for scientific inquiries

Contact

Name of organization / entity

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Full name of responsible person

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Position

Gynecologist

Other areas of specialty/work

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Person responsible for updating data

Contact

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Position

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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