

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

01 Jul 2026

### Comparison of efficacy of vitamin E and alpha lipoic acid in decrease of oxidative stress in children with Down syndrome

#### Protocol summary

##### Summary

This study is a Randomized Controlled Trial to evaluate the effects of dietary alpha tocopherol and alpha lipoic acid supplements for reducing oxidative stress in children with Down syndrome. The study group consists of 105 children with DS aged 7-15 years. All children will select from Special Education Centers for Mentally Handicapped Children in different districts of Tehran, Iran. After selection of the centres parents will be invited to attend an information session. We will describe the procedures and details of the trial and will ask them to sign an informed consent form if they desire to participate their children in the trial. After that we allocate them to three groups randomly: alpha tocopherol (400 IU/d), ALA (100 mg/d) and placebo. The intervention period will take four months. Serum malondialdehyde (MDA) and urinary 8-hydroxydeoxyguanosine (8OHdG) will use as biomarkers of oxidative stress. These biomarkers will be measured on before and after intervention. For this reason we will take 5cc blood and 10 cc urine samples from children at baseline prior to supplementation and after 4 months of supplementation.

#### General information

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT2013102215111N1**

Registration date: **2013-10-30, 1392/08/08**

Registration timing: **retrospective**

Last update:

Update count: **0**

##### Registration date

2013-10-30, 1392/08/08

##### Registrant information

**Name**

Seyyed Mostafa Nachvak

##### Name of organization / entity

kermanshah university of medical sciences

##### Country

Iran (Islamic Republic of)

##### Phone

+98 83 1826 2005

##### Email address

m.nachrak@kums.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

Exceptional children research

##### Expected recruitment start date

2006-01-21, 1384/11/01

##### Expected recruitment end date

2006-05-22, 1385/03/01

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

##### Scientific title

Comparison of efficacy of vitamin E and alpha lipoic acid in decrease of oxidative stress in children with Down syndrome

##### Public title

The effect of dietary supplements of alpha lipoic acid and vitamin E on stress oxidative in children with Down syndrome

##### Purpose

Treatment

##### Inclusion/Exclusion criteria

Inclusion criteria: confirmation of regular trisomy 21 in children by cytogenetic analysis; not on any medication or vitamin supplementation therapy for six months before the start of intervention.

## Age

From **7 years** old to **15 years** old

## Gender

Both

## Phase

N/A

## Groups that have been masked

*No information*

## Sample size

Target sample size: **105**

## Randomization (investigator's opinion)

Randomized

## Randomization description

## Blinding (investigator's opinion)

Double blinded

## Blinding description

## Placebo

Used

## Assignment

Parallel

## Other design features

## Secondary Ids

empty

## Ethics committees

### 1

#### Ethics committee

##### Name of ethics committee

Tehran university of medical sciences

##### Street address

Tehran - Porsina St

##### City

Tehran

##### Postal code

#### Approval date

2005-12-27, 1384/10/06

#### Ethics committee reference number

132/8316

## Health conditions studied

### 1

#### Description of health condition studied

Down syndrome

#### ICD-10 code

Q90

#### ICD-10 code description

Down syndrome

## Primary outcomes

### 1

#### Description

MDA

#### Timepoint

before intervention and after 4 months

#### Method of measurement

The determination of MDA levels will performed by method of Satho . In this method the reaction of MDA with Thiobarbituric acid (TBA) creates a complex which is determined spectrophotometrically.

### 2

#### Description

urinary 8-hydroxydeoxyguanosine (8OHdG)

#### Timepoint

before intervention and after 4 months

#### Method of measurement

8OHdG in urine samples will assessed using an enzyme-linked immunosorbent assay (ELISA) kit (8OHdG Quantitation, Cell Biolabs, Inc. San Diego ).

## Secondary outcomes

empty

## Intervention groups

### 1

#### Description

In group 1 a capsule contained 400 IU alpha tocopherol will give to the children every day after dinner or lunch meal for 4 months.

#### Category

Treatment - Other

### 2

#### Description

In group 2 a capsule contained 100 mg alpha lipoic acid will give to the children every day after dinner or lunch meal for 4 months.

#### Category

Treatment - Drugs

### 3

#### Description

In group 3 a capsule contained only vegetable oil as a placebotocopherol will give to the children every day after dinner or lunch meal for 4 months.

#### Category

Treatment - Drugs

## Recruitment centers

### 1

#### Recruitment center

##### Name of recruitment center

Department of exceptional children of Tehran

##### Full name of responsible person

Dr Azam Hoseinipor

##### Street address

Tehran Blv kashani

##### City

## Sponsors / Funding sources

1

### Sponsor

**Name of organization / entity**

Exceptional Children Institute

**Full name of responsible person**

Dr. Kazem Dastjerdi

**Street address**

Gharani No 23

**City**

Tehran

**Grant name**

تحقیقاتی

**Grant code / Reference number**

0259

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

Yes

**Title of funding source**

Exceptional Children Institute

**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**

Exceptional Children Research

**Full name of responsible person**

Dr Seyyed Mostafa Nachvak

**Position**

Academic member/PhD on Nutrition

**Other areas of specialty/work**

**Street address**

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## Person responsible for scientific inquiries

### Contact

**Name of organization / entity**

Tehran University of Medical Sciences

**Full name of responsible person**

Dr Seyyed Ali Keshavarz

**Position**

Full professor/PhD of Nutrition

**Other areas of specialty/work**

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## Person responsible for updating data

### Contact

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**Full name of responsible person**

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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*