

Clinical Trial Protocol

Iranian Registry of Clinical Trials

13 Jun 2026

Study of zinc supplement on oxidative stress and inflammation markers in multiple myeloma patients undergo autologous hematopoietic stem cell transplantation compared with placebo

Protocol summary

Study aim

Determination of the level of markers and genes of oxidative stress and inflammation in patients undergoing autologous BMT under the influence of oral zinc supplementation in comparison with the placebo group

Design

This study is a double-blind, placebo-controlled clinical trial that will be randomized by using a block design method, and study participants are randomly divided into two groups of 20 people, Zinc and Placebo. In the first month after transplantation, the patients will receive three tablets of zinc gluconate or placebo. Zinc and copper Serum levels will be measured before intervention and on days 15 and 30 after HSCT. The expression of Nrf2, INrf2 (Keap1), SOD1, SOD2, NOX1, NOX2 genes as well as the levels of TNF- α and the activity of MDA, NO enzymes will be measured to determine the effect of oral zinc supplementation on gene expression, enzyme levels and measure inflammatory and oxidative stress markers.

Settings and conduct

Eligible patients will be selected from those referred to the BMT department of Taleghani Hospital for auto-HSCT. Patients enter the study after describing the objectives of the study and filling out the consent form.

Participants/Inclusion and exclusion criteria

Inclusion Criteria: History of Multiple Myeloma; A complete response to treatment; Patients are candidates for AHST; have a health certificate. Exclusion Criteria: History of sensitivity to oral zinc supplement; The patients who have consumed oral zinc supplementation since three months before the transplantation; Zinc serum levels above 200mg/dl.

Intervention groups

Group 1: Patients with a history of multiple myeloma undergoing BMT who receive placebo. Group 2: Patients with a history of multiple myeloma undergoing BMT who

receive zinc.

Main outcome variables

Expression of Nrf2, INrf2 (Keap1), SOD1, SOD2, NOX1, NOX2 genes; TNF- α cytokine; Activity of MDA, NO enzymes.

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20211227053546N1**

Registration date: **2022-03-23, 1401/01/03**

Registration timing: **registered_while_recruiting**

Last update: **2022-03-23, 1401/01/03**

Update count: **0**

Registration date

2022-03-23, 1401/01/03

Registrant information

Name

Nariman Mosaffa

Name of organization / entity

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2022-03-21, 1401/01/01

Expected recruitment end date

2022-09-22, 1401/06/31

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Study of zinc supplement on oxidative stress and inflammation markers in multiple myeloma patients undergo autologous hematopoietic stem cell transplantation compared with placebo

Public title

Zinc supplement effect on oxidative stress and inflammation markers in multiple myeloma patients undergo autologous hematopoietic stem cell transplantation

Purpose

Treatment

Inclusion/Exclusion criteria**Inclusion criteria:**

Complete response to treatment Lack of underlying disease Do not get any infection in the last three months, including COVID19 Age range 40 to 60 years Patients with a history of multiple myeloma

Exclusion criteria:

History of sensitivity to "zinc" oral supplement Serum levels above 200 mg/dl Patients who have taken oral supplements for three months before transplantation. People's unwillingness to collaborate in the study

Age

From **40 years** old to **60 years** old

Gender

Both

Phase

2-3

Groups that have been masked

- Participant
- Investigator

Sample size

Target sample size: **40**

Randomization (investigator's opinion)

Randomized

Randomization description

In this randomization method, the size of the blocks is considered equal. The size of each block is 4, which includes 2 participants in the zinc group and 2 participants in the placebo group. The SNOSE method is used to perform randomization. Based on the sample size of the research, a number of envelopes were prepared with aluminum wrappers and each of the random sequences created on a card was recorded and the cards were placed in the envelopes of the letter, respectively. In order to maintain a random sequence, the envelopes were numbered in the same way on the outer surface. Finally, the lids of the letter envelopes were glued and placed in a box, respectively. At the beginning of the registration of the participants, according to the order of entry of the eligible participants, one of the envelopes will be opened in order

and the assigned group of that participant will be revealed. Random sequencing was performed by SAS software version 9.4

Blinding (investigator's opinion)

Double blinded

Blinding description

This clinical trial is performed in a double-blind method, which means the patient and the principal researcher are not aware of the patient's treatment. Therefore, to maintain blindness, the randomization list is not shown to the principal researcher. A statistician prepares the randomization list. The package evaluator provides a form containing either zinc or a placebo for each patient. Zinc and placebo tablets are the same color and shape but have different ingredients.

Placebo

Used

Assignment

Parallel

Other design features

In this study, based on similar studies and in consultation with its researchers, the amount of "zinc" in 30 days after transplantation was selected to be 90 mg daily

Secondary Ids

empty

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

Ethics committee of Shahid beheshti University of Medical Sciences_ Medical school

Street address

Tehran - Shahid Chamran Highway Shahid Shahriari Square - Koodkiar Street - School of Medicine

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Province

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Postal code

1985717443

Approval date

2022-02-09, 1400/11/20

Ethics committee reference number

IR.SBMU.MSP.REC.1400.728

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

Patients with multiple myeloma who are candidates for stem cell transplantation.

ICD-10 code

C90

ICD-10 code description

Multiple myeloma and malignant plasma cell neoplasms

Primary outcomes

1

Description

Determination of genes(Nrf2-INrf2(Keap1)-SOD1,2-NOX1,2) expression

Timepoint

Before hematopoietic stem cell transplantation beginning and 15,30 days after transplantation

Method of measurement

In DNA extracted from peripheral blood mononuclear cells using Real-Time-PCR

2

Description

Determination of MDA-NO protein levels

Timepoint

Before hematopoietic stem cell transplantation beginning and 15,30 days after transplantation

Method of measurement

MDA assay by thiobarbituric acid method and NO assay by griess method

3

Description

Determination of TNF-a cytokine

Timepoint

Before hematopoietic stem cell transplantation beginning and 15,30 days after transplantation

Method of measurement

Assay of cytokines by ELISA method

Secondary outcomes

empty

Intervention groups

1

Description

Intervention group: Patients who underwent autologous HSCT receive three tablets of "gluconate zinc" (each tablet containing 30 mg zinc elemental) daily after eating main meals for first 30 days post-HSCT

Category

Treatment - Drugs

2

Description

Control group: Patients who underwent autologous HSCT receive three tablets of "placebo" (similar to zinc supplements) daily after eating the main meals for first 30 days post-HSCT

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Ayatollah Taleghani Educational Hospital, Shahid Beheshti University of Medical Sciences

Full name of responsible person

Abbas Hajifathali

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Arabian Street, End of Velenjak Street, Shahid Chamran Highway

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

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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

Shahid Beheshti 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**

Shahid Beheshti University of Medical Sciences

Full name of responsible person

Nariman Mosaffa

Position

Professor

Latest degree

Ph.D.

Other areas of specialty/work

Immunology

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Person responsible for scientific inquiries**Contact****Name of organization / entity**

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Person responsible for updating data**Contact****Name of organization / entity**

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

No - There is not a plan to make this available

Justification/reason for indecision/not sharing IPD

For ethical reasons, we have no plans to publish
participants' data

Study Protocol

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

Statistical Analysis Plan

Not applicable

Informed Consent Form

No - There is not 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

No - There is not a plan to make this available

Data Dictionary

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