

Clinical Trial Protocol

Iranian Registry of Clinical Trials

17 Sep 2026

Effect of vitamin D, omega 3 fatty acids supplementation and co-supplementation of them on liver, inflammatory biomarkers, and oxidative stress in patients with colorectal cancer

Protocol summary

Study aim

Evaluation of the effect of vitamin D, omega 3 fatty acids supplementation and co-supplementation of them on liver, inflammatory biomarkers, and oxidative stress in patients with colorectal cancer

Design

The study is done in Phase 3 clinical trials with a control group, in a factorial design, double-blind and randomized allocation. 80 eligible colorectal cancer patients were randomly allocated into 4 groups (vitamin D, omega-3, co-supplementation, placebo). For randomized allocation performing, permuted block randomization will be used by quadrilateral blocks.

Settings and conduct

The participants, in Oncology clinic of Tehran Gastroenterology & Hepatology Center, who meet the criteria, will be randomly allocated into 4 groups. Blood samples were taken to quantify serum levels of 25(OH)D, and liver enzymes at baseline and after 8 weeks of intervention.

Participants/Inclusion and exclusion criteria

Inclusion criteria: Patients with stage II or III colorectal cancer and were candidate to receive chemotherapy; Aged 18 years or more; BMI range of 18.5-30 kg/m²; Serum 25(OH)D < 30 ng/ml; Not having autoimmune or chronic diseases; No taking vitamin D and/or omega-3 supplements and other vitamin- mineral supplements; Not taking laxative and anti-inflammatory medications; Not allergic to fish and fish products.

Intervention groups

The intervention groups, for an 8-week period, will be randomly allocated into 4 groups: 1) a 50000 IU vitamin D tablet (Zahravi, Iran), weekly+ 2 omega-3 fatty acid capsules (each capsule containing 330 mg omega-3 fatty acid), daily (Minami Nutrition, Belgium); 2) a 50000 IU vitamin D tablet, weekly+ 2 omega-3 fatty acid placebo (Zahravi, Iran), daily; 3) a vitamin D placebo (Zahravi,

Iran), weekly+ 2 omega-3 fatty acid capsules; 4) a vitamin D placebo, weekly+ 2 omega-3 fatty acid placebo, daily.

Main outcome variables

Serum level of TNF- α

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20090822002365N22**

Registration date: **2018-12-30, 1397/10/09**

Registration timing: **prospective**

Last update: **2018-12-30, 1397/10/09**

Update count: **0**

Registration date

2018-12-30, 1397/10/09

Registrant information

Name

Mohammad Reza Vafa

Name of organization / entity

Iran University of Medical Sciences

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2019-01-05, 1397/10/15

Expected recruitment end date

2019-07-06, 1398/04/15

Actual recruitment start date
empty

Actual recruitment end date
empty

Trial completion date
empty

Scientific title
Effect of vitamin D, omega 3 fatty acids supplementation and co-supplementation of them on liver, inflammatory biomarkers, and oxidative stress in patients with colorectal cancer

Public title
Effect of vitamin D and omega 3 fatty acids co-supplementation on colorectal cancer

Purpose
Treatment

Inclusion/Exclusion criteria
Inclusion criteria:
Patients with stage II or III colorectal cancer and were candidate to receive chemotherapy; aged 18 years or more; body mass index range of 18.5-30 kg/m²; serum 25(OH)D < 30 ng/ml; not having autoimmune or chronic diseases; not taking vitamin D and/or omega-3 supplements and other vitamin- mineral supplements or parenteral nutrition; not taking laxative and anti-inflammatory medications; not allergic to fish and fish products;
Exclusion criteria:

Age
From 18 years old

Gender
Both

Phase
3

Groups that have been masked

- Participant
- Care provider
- Investigator

Sample size
Target sample size: 80

Randomization (investigator's opinion)
Randomized

Randomization description
For randomized allocation performing, permuted block randomization will be used by quadrilateral blocks. According to the sample size of 80 subjects, 20 blocks will be generated using the online site (www.sealedenvelope.com). In order to allocation concealment in the randomized process, unique codes will be used on the drug boxes that is generated by the software. Participants will enter based on the sequence produced into study and the drug packets with code registered will allocate to the individual. Therefore, before participants selection, they will be unaware of the type of intervention that will receive, as well as a random sequence produced during the study will be immune from prediction.

Blinding (investigator's opinion)

Double blinded

Blinding description
For blinding, a person who is not involved in study protocol created the randomization list assigning participants to the vitamin D, omega-3, co-supplementation or the placebo group. Vitamin D, omega-3, and placebo tablets are placed into unlabeled identical containers. The study leader will label these containers with participant numbers using the randomization list. All investigators, and participants were blinded to the random assignments.

Placebo
Used

Assignment
Factorial

Other design features

Secondary Ids
empty

Ethics committees

1

Ethics committee
Name of ethics committee
Ethics committee of Iran University of Medical of Sciences
Street address
Iran University of Medical Science, Hemmat Highway
City
Tehran
Province
Tehran
Postal code
1449614535

Approval date
2018-12-13, 1397/09/22

Ethics committee reference number
IR.IUMS.REC.1397.572

Health conditions studied

1

Description of health condition studied
Colorectal cancer

ICD-10 code
C18.9

ICD-10 code description
Malignant neoplasm of colon, unspecified

Primary outcomes

1

Description
Serum level of TNF- α .

Timepoint
Before and 8 weeks after intervention

Method of measurement

Serum TNF- α levels will be assessed by ELISA and Bender Med kit (Bender Med, Germany).

Secondary outcomes

1

Description

Serum levels of Matrix Metalloproteinase-2 (MMP-2)

Timepoint

Before and 8 weeks after intervention.

Method of measurement

Serum levels of MMP-2 will be assessed by ELISA and Zellbio kit (Zellbio, Germany).

2

Description

Serum levels of Matrix Metalloproteinase-9 (MMP-9)

Timepoint

Before and 8 weeks after intervention.

Method of measurement

Serum levels of MMP-9 will be assessed by ELISA and Zellbio kit (Zellbio, Germany).

3

Description

Serum levels of Alkaline phosphatase (ALP)

Timepoint

Before and 8 weeks after intervention.

Method of measurement

Serum levels of ALP will be assessed by ELISA and Zellbio kit (Zellbio, Germany).

Intervention groups

1

Description

Intervention 1: a 50000 IU vitamin D tablet (Zahravi, Iran), weekly+ 2 omega-3 fatty acid capsules (each capsule containing 330 mg omega-3 fatty acid) (Minami Nutrition, Belgium), daily, for 8 weeks.

Category

Treatment - Drugs

2

Description

Intervention group 2: a 50000 IU vitamin D tablet (Zahravi, Iran), weekly+ 2 omega-3 fatty acid placebo (Zahravi, Iran), daily, for 8 weeks.

Category

Treatment - Drugs

3

Description

Intervention group 3: two omega-3 fatty acid capsules (each capsule containing 330 mg omega-3 fatty acid) (Minami Nutrition, Belgium), daily+ a vitamin D placebo

(Zahravi, Iran), weekly, for 8 weeks.

Category

Treatment - Drugs

4

Description

Control group: a vitamin D placebo (Zahravi, Iran), weekly+ 2 omega-3 fatty acid placebo (Zahravi, Iran), daily, for 8 weeks.

Category

Placebo

Recruitment centers

1

Recruitment center

Name of recruitment center

Oncology clinic of Tehran Gastroenterology & Hepatology Center (TGHC)

Full name of responsible person

Dr. Masood Iravani

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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
Iran 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
Iran University of Medical Sciences
Full name of responsible person
Dr. Mohammadreza Vafa
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Professor
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Sharing plan

Deidentified Individual Participant Data Set (IPD)

Yes - There is a plan to make this available

Study Protocol

Yes - There is a plan to make this available

Statistical Analysis Plan

Yes - There is a plan to make this available

Informed Consent Form

Yes - There is a plan to make this available

Clinical Study Report

Yes - There is a plan to make this available

Analytic Code

Yes - There is a plan to make this available

Data Dictionary

Yes - There is a plan to make this available

Title and more details about the data/document

A part of the data will be shared, such as primary outcomes.

When the data will become available and for how long

4 months after the publication of the results.

To whom data/document is available

researchers and students of university.

Under which criteria data/document could be used

Four months after the publication of this study papers, the obtained data will be available to the applicant researchers and students for further analysis.

From where data/document is obtainable

Applicants can be contacted with corresponding author by e-mail or postal address to receive the requested data. Postal address: Nutrition Department, School of health, Iran University of Medical Science, Hemmat Expressway, Tehran Phone Number: 0098 2186704743

E-mail: rezavafa@yahoo.com

What processes are involved for a request to access data/document

Applicants can be contacted with corresponding author by e-mail or postal address to receive the requested data. Postal address: Nutrition Department, School of health, Iran University of Medical Science, Hemmat Expressway, Tehran Phone Number: 0098 2186704743
E-mail: rezavafa@yahoo.com

Comments