

Clinical Trial Protocol

Iranian Registry of Clinical Trials

06 Jun 2026

Effect of spearmint -ginger capsule on nausea and vomiting in patients undergoing chemotherapy for breast cancer in Tohid hospital in Sanandaj, 2018.

Protocol summary

Study aim

Determination of the effect of Peppermint Capsule - Ginger on Nausea and Vomiting of Chemotherapy Patients with Breast Cancer in Sanandaj Tohid Medical Center

Design

Randomized Clinical trial with control group; parallel groups, blinded, sample size 70

Settings and conduct

Clinical, triple blind clinical trials (chemotherapy patients; research collaborator and counselor) will be conducted at Tohid Hospital in Sanandaj.

Participants/Inclusion and exclusion criteria

Entry requirements: Patients are females and over 18 years old; At least the reading and writing literacy of the patient or one of the main members of the patient's family; A definitive diagnosis of breast cancer by an oncologist and pathologist;. History of at least one chemotherapy course; History of nausea and vomiting after previous chemotherapy sessions; Absence of simultaneous radiotherapy with chemotherapy; Do not take warfarin, heparin and aspirin therapy; No history of blood disorders such as thrombocytopenia and leukopenia; Not having a history of allergy to ginger or peppermint and their use during the last week

Intervention groups

In the intervention group, a 500 mg ginger mixture containing 250 mg of ginger and 250 mg of peppermint powder will be given to chemotherapy patients. The intervention will take place about five hours before the chemotherapy completion. During this period, along with the usual anti-nausea regimen (dexamethasone ampoules, cholesterol ampoules and apoptotic pills capsules), each day, two 500 mg ginger-mint capsules will be consumed at a time of 12 hours (a total of 1 g), preferably with an empty stomach. In the control group, a 500 mg starch capsule will be used.

Main outcome variables

Incidence of nausea, Severity of nausea, Incidence of vomiting

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20180215038741N1**

Registration date: **2018-04-23, 1397/02/03**

Registration timing: **prospective**

Last update: **2018-04-23, 1397/02/03**

Update count: **0**

Registration date

2018-04-23, 1397/02/03

Registrant information

Name

Havre Mawloudi

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 87 3633 5014

Email address

Mawloudi.h@muk.ac.ir

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2018-05-04, 1397/02/14

Expected recruitment end date

2018-08-05, 1397/05/14

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Effect of spearmint -ginger capsule on nausea and vomiting in patients undergoing chemotherapy for breast cancer in Tohid hospital in Sanandaj, 2018.

Public title

Effect of spearmint -ginger capsule on nausea and vomiting

Purpose

Treatment

Inclusion/Exclusion criteria**Inclusion criteria:**

History of at least one course of chemotherapy History of nausea and vomiting after previous chemotherapy sessions Definitive diagnosis of breast cancer by clinical trials and an oncologist

Exclusion criteria:

Having a history of allergy to ginger or mint and taking them over the past week Having other nausea-causing problems such as high blood pressure, liver failure, kidney problems and digestive problems Metastasis to other organs of other body systems

Age

From **18 years** old

Gender

Female

Phase

1-2

Groups that have been masked

- Participant
- Care provider
- Data analyser

Sample size

Target sample size: **70**

Randomization (investigator's opinion)

Randomized

Randomization description

In this study, by using the available sampling method and the random identification method, by using the referral list, the samples will first be selected based on entry criteria. Then, each eligible individuals will be allocated into control or intervention group.

Blinding (investigator's opinion)

Triple blinded

Blinding description

Patients undergoing chemotherapy; Research collaborators and counselors from the intervention and control group will be unaware.

Placebo

Used

Assignment

Parallel

Other design features**Secondary Ids**

empty

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

Ethics Committee of Kurdistan University of Medical Sciences

Street address

Kurdistan University of Medical Sciences, after Qods Hospital, Sanandaj

City

Sanandaj

Province

Kurdistan

Postal code

66617713446

Approval date

2018-03-24, 1397/01/04

Ethics committee reference number

IR.MUK.REC.1396/370

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

Breast cancer patients

ICD-10 code**ICD-10 code description****Primary outcomes****1****Description**

Incidence of nausea, Severity of nausea, Incidence of vomiting

Timepoint

An hour before chemotherapy, during chemotherapy, first, second, third, fourth and fifth days of chemotherapy

Method of measurement

A questionnaire, an information recording form, and a standard analogue visual analogue tool

Secondary outcomes**1****Description****Timepoint****Method of measurement****Intervention groups**

1

Description

Intervention group: combination of spearmint _ Ginger capsules 500 mg containing 250 mg of ginger and 250 mg spearmint powder, every day two capsules will be given to chemotherapy patients within 12 hours. The intervention will begin one hour before the chemotherapy and will last for five days. Training will be given to patients in the early hours of chemotherapy. The capsules will be purchased by Barij Essen, Isfahan, with the standard code C200464 to the 1028 registration number.

Category

Treatment - Drugs

2

Description

Control group: Starch 500 mg capsules every day two capsules will be given to chemotherapy patients within 12 hours. The intervention will begin an hour before the chemotherapy and will last for five days. Training will be given to patients in the early hours of chemotherapy. The capsules will be purchased by Barij Essen, Isfahan, with the standard code C200464 to the 1028 registration number

Category

Placebo

Recruitment centers

1

Recruitment center

Name of recruitment center

Tohid Hospital

Full name of responsible person

Jamal Seidi

Street address

Kurdistan University of Medical Sciences, after Qods Hospital, Sanandaj sanandaj Kurdistan Iran

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Mawloudi.h@muk.ac.ir

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Sanandaj University of Medical Sciences

Full name of responsible person

Student Research Committee, Vice Chancellor for

research and technology of Kurdistan University of M

Street address

Kurdistan University of Medical Sciences, after Qods Hospital, Pasdaran street, Sanandaj sanandaj kurdistan Iran, Islamic Republic Of

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Web page address

Grant name

Grant code / Reference number

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

Yes

Title of funding source

Sanandaj 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

Sanandaj University of Medical Sciences

Full name of responsible person

Jamal Seidi

Position

ph.D/Assistant professor

Latest degree

Ph.D.

Other areas of specialty/work

Nursery

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Person responsible for scientific inquiries

Contact

Name of organization / entity

Sanandaj University of Medical Sciences

Full name of responsible person

Jamal Seidi

Position

ph.D/Assistant professor

Latest degree

Ph.D.

Other areas of specialty/work

Nursery

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Person responsible for updating data

Contact

Name of organization / entity

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Full name of responsible person

havre mawloudi

Position

Senior Surgical Student

Latest degree

Master

Other areas of specialty/work

Nursery

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Email

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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