

Clinical Trial Protocol

Iranian Registry of Clinical Trials

31 May 2026

Comparison the effects of aromatherapy with peppermint oil and routine drug for chemotherapy induced nausea and vomiting in patients with breast cancer

Protocol summary

Summary

The objective of this study is to determine the effect of Aromatherapy with peppermint oil on chemotherapy-induced nausea and vomiting in patients with Breast cancer. This study is a Randomized clinical trial. The study population included 100 women who referred to Rasoul Akram hospital and Cancer Institute of Iran. Inclusion criteria: patients Aged 18-65 years, who Receiving cisplatin and anthracycline for the first cycle of chemotherapy. Patients who are reluctant to complete the study or have allergy to peppermint oil will be excluded. After obtaining inform consent, convenience sampling method will be used to recruit the study subjects, then participants will randomly allocated to the intervention group and the control group. The intervention group, besides the standard medications prescribed by a doctor, will trained to pour 2 drops pure peppermint oil (Barij Essence Kashan) by dropper on a napkin (size 20 × 20 cm) and attach with a pin on their collar then will be asked for taking breathe normally for 20 minutes. They will teach to do it three times each day (morning, noon and evening) for five consecutive days. For control group will be used normal saline as a placebo. To assess the nausea and vomiting, Rhodes questionnaire will be completed for the subsequent 5 days by both the intervention and the control group.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2014050215649N2**

Registration date: **2014-11-14, 1393/08/23**

Registration timing: **registered_while_recruiting**

Last update:

Update count: **0**

Registration date

2014-11-14, 1393/08/23

Registrant information

Name

Mohammad Eghbali

Name of organization / entity

Torbat Heydariyeh University of Medical Sciences

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

School of Nursing and Midwifery, Tehran University of Medical Sciences

Expected recruitment start date

2014-11-01, 1393/08/10

Expected recruitment end date

2015-04-30, 1394/02/10

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Comparison the effects of aromatherapy with peppermint oil and routine drug for chemotherapy induced nausea and vomiting in patients with breast cancer

Public title

The effects of aromatherapy with peppermint oil for chemotherapy induced nausea and vomiting

Purpose

Prevention

Inclusion/Exclusion criteria

Inclusion criteria: patients Aged 18-65 years, who Receiving cisplatin and anthracycline for the first cycle of chemotherapy, Healthy sense of smell. Exclusion criteria: reluctant to complete the study or have allergy to peppermint oil.

Age

From **18 years** old to **65 years** old

Gender

Female

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **100**

Randomization (investigator's opinion)

Randomized

Randomization description

Blinding (investigator's opinion)

Not blinded

Blinding description

Placebo

Used

Assignment

Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethics committee of Tehran University of Medical Science

Street address

Sixth Floor, Central Organization of University, Qods St, Keshavarz Blvd

City

Tehran

Postal code

Approval date

2014-05-18, 1393/02/28

Ethics committee reference number

93/02/28/25630

Health conditions studied

1

Description of health condition studied

Nausea and vomiting

ICD-10 code

R11

ICD-10 code description

Nausea and vomiting

Primary outcomes

1

Description

Acute and delayed nausea

Timepoint

Data are gathered for five days after starting chemotherapy

Method of measurement

Rhodes Questionnaire

Secondary outcomes

1

Description

Acute and delayed vomiting

Timepoint

Data are gathered for five days after starting chemotherapy

Method of measurement

Rhodes Questionnaire

Intervention groups

1

Description

The intervention group, besides the standard medications prescribed by a doctor, will be trained to pour 2 drops pure peppermint oil (Barij Essence Kashan) by dropper on a napkin (size 20 × 20 cm) and attach with a pin on their collar then will be asked for taking breathe normally for 20 minutes. They will teach to do it three times each day (morning, noon and evening) for five consecutive days

Category

Prevention

2

Description

For control group will be used normal saline as a placebo.

Category

Placebo

Recruitment centers

1

Recruitment center

Name of recruitment center

Rasoul akram hospital

Full name of responsible person

Mohammad Eghbali

Street address

Niayesh St, sattarkhan St.

City
Tehran

2

Recruitment center

Name of recruitment center
Cancer Institute of Imam Khomeini Hospital in Tehran
Full name of responsible person
Mohammad Eghbali
Street address
Cancer Institute Central 2, Dr. Gharib St.
City
Tehran

Sponsors / Funding sources

1

Sponsor

Name of organization / entity
Vice chancellor for research, Tehran University of
Medical Sciences
Full name of responsible person
Doctor Masood Younesian - Vice President of
Research and Technology
Street address
Keshavarz St, Qods St., Central Building, Sixth Floor,
Tehran University of Medical Sciences, Tehran
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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, Tehran 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
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Full name of responsible person
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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