

Clinical Trial Protocol

Iranian Registry of Clinical Trials

22 Jun 2026

The effect of oral rinse Peppermint Essence (PE) in prevention of chemotherapy- induced oral mucositis

Protocol summary

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Summary

Purpose: This study has been conducted to determine the effect of using oral rinse Peppermint Essence (PE) in prevention of chemotherapy-induced oral mucositis (OM). Materials & Methods: 40 patients with colon & rectum cancer, who admitted for chemotherapeutic management, were enrolled in a double blind clinical trial and they randomly divided in two equal, placebo & PE groups. They had to receive 10 drop of oral rinse PE or placebo, tree times a day, from the first day of chemotherapy up to the fourteenth day. Data tools included a bipartite questionnaire (demographic Q & clinical Q) and World Health Organization (WHO) scale for measurement of OM.

Recruitment status

Recruitment complete

Funding source

Private

Expected recruitment start date

2009-09-23, 1388/07/01

Expected recruitment end date

2009-12-11, 1388/09/20

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT138809222846N1**

Registration date: **2010-05-19, 1389/02/29**

Registration timing: **retrospective**

Last update:

Update count: **0**

Registration date

2010-05-19, 1389/02/29

Registrant information

Name

Tahereh Ashktorab

Name of organization / entity

Shahid Beheshti University of Medical Sciences

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Iran (Islamic Republic of)

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Email address

Scientific title

The effect of oral rinse Peppermint Essence (PE) in prevention of chemotherapy- induced oral mucositis

Public title

The effect of oral rinse Peppermint Essence (PE) in prevention of chemotherapy- induced oral mucositis

Purpose

Prevention

Inclusion/Exclusion criteria

Inclusion criteria: consciousness/literate/ admitted to hospital with colon or rectum cancer/ chemotherapeutic management with 5-fluorouracil / no stomatitis/ no oral disease/ no asthma/ no renal & liver disorders/ no auto immune disease and diabetes/ no fever and neutropenia/ no recieved antibiotic & narcotic analgesic/ no recieved another oral rinse /no mint family allergy.

Age

From **30 years** old to **70 years** old

Gender

Both

Phase

2-3

Groups that have been masked

No information

Sample size

Target sample size: 40

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

The International Branch of Shahid Beheshti
University of Medical Sciences

Street address

No. 19, Shaheed Abbas pour, Vali Asr

City

Tehran

Postal code

14155-3693

Approval date

2009-09-20, 1388/06/29

Ethics committee reference number

116/677

Health conditions studied

1

Description of health condition studied

oral mucositis

ICD-10 code

K13

ICD-10 code description

Other diseases of lip and oral mucosa

Primary outcomes

1

Description

oral mucositis

Timepoint

daily

Method of measurement

observation with a check list

Secondary outcomes

empty

Intervention groups

1

Description

Intervention: 10 drops of oral rinse Peppermint Essence,
tree times a day, from the first day of chemotherapy up
to the fourteenth day

Category

Prevention

2

Description

Control: placebo

Category

Placebo

Recruitment centers

1

Recruitment center

Name of recruitment center

Tajrish hospital

Full name of responsible person

Street address

City

Tehran

2

Recruitment center

Name of recruitment center

Emam Hosein hospital

Full name of responsible person

Street address

City

Tehran

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Researcher

Full name of responsible person

Tahereh Ashktorab

Street address

Faculty of Nursing, Shahid Beheshti University of
Medical Sciences

City

Tehran

Grant name

Grant code / Reference number

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

Yes
Title of funding source
Researcher
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

PhD
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Person responsible for general inquiries

Contact

Name of organization / entity
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Person responsible for updating data

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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