

Clinical Trial Protocol

Iranian Registry of Clinical Trials

09 Jun 2026

Effect of inhalation of ginger essence on nausea and vomiting after nephrectomy

Protocol summary

Summary

In this randomized clinical study, single blind for experimental groups, 120 eligible patient will randomly be allocated between the intervention and the control group. After surgical procedure, When patients are transferred to the recovery room, in the intervention group 2 drops of ginger essence will be dripped on A2*2 gaze and this will be repeated every 15 minute during 2 hours of patients stay in recovery room. Nausea and vomiting will be checked every 15 minute for 2hours and then after 6 hours when the patients are transferred to ward. In the control group the same procedure will be done but 2 drops of normal saline will be used instead of ginger. Nausea will be assessed using a Visual analog scale.

General information

Acronym

Effect of ginger essence on PONV

IRCT registration information

IRCT registration number: **IRCT2014080218650N1**

Registration date: **2014-09-07, 1393/06/16**

Registration timing: **registered_while_recruiting**

Last update:

Update count: **0**

Registration date

2014-09-07, 1393/06/16

Registrant information

Name

Fatemeh Sadat Hosaini

Name of organization / entity

Kashan University of Medical Sciences

Country

Iran (Islamic Republic of)

Phone

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Email address

fatemehhosaini57@kaums.ac.ir

Recruitment status

Recruitment complete

Funding source

Vice Chancellor for research of Kashan University of Medical Sciences

Expected recruitment start date

2014-09-01, 1393/06/10

Expected recruitment end date

2015-01-02, 1393/10/12

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Effect of inhalation of ginger essence on nausea and vomiting after nephrectomy

Public title

Effect of inhalation of ginger essence on nausea and vomiting after nephrectomy

Purpose

Prevention

Inclusion/Exclusion criteria

Inclusion criteria: age of 18 years or older; any known allergy to ginger; not having motion sickness (car-sea); not receiving anti-emetic drugs; not receiving warfarin, heparin, aspirin 325 mg; no receiving chemotherapy drugs; being NPO for at least 8 hours before surgery, not having a history of asthma and other respiratory disorder; not having an olfactory sensation disorders, ability to read and write; not having disorders of brain stem and cerebellum, not having a coagulopathy. Exclusion criteria: occurring any allergic reaction to ginger during the study; intolerance to ginger essence

odor.
Age
No age limit
Gender
Both

Phase
N/A
Groups that have been masked
No information
Sample size
Target sample size: **120**
Randomization (investigator's opinion)
Randomized
Randomization description
Blinding (investigator's opinion)
Single blinded
Blinding description
Placebo
Used
Assignment
Parallel
Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee
Name of ethics committee
" Ethics committee Kashan University of Medical Sciences"
Street address
Ghotbe ravandi road, km 5.
City
Kashan
Postal code
8715981151
Approval date
2014-07-22, 1393/04/31
Ethics committee reference number
29/5/1/1899/p

Health conditions studied

1

Description of health condition studied
Nausea and Vomiting
ICD-10 code
R11
ICD-10 code description
Other diseases of the digestive system

2

Description of health condition studied
kidney surgery
ICD-10 code

XIV
ICD-10 code description
Other disorders of kidney

Primary outcomes

1

Description
Nausea
Timepoint
Every 15 minute during surgery and 2 hours and 6 hours after surgery.
Method of measurement
Visual Analog Scale

2

Description
Vomiting
Timepoint
number in the first 2 hours after the nephrectomy and after 6 hours
Method of measurement
Counting the number of vomiting

Secondary outcomes

empty

Intervention groups

1

Description
Intervention: 2 drops of 100% ginger essence will be used on a piece of gauze that will be attached to the collar of the patients clothes every 15 minute for 2 hour in the recovery room.
Category
Treatment - Drugs

2

Description
control:normal saline,2 drop normal saline 9% every 15 minute for 2 hour to recovery room
Category
Treatment - Drugs

Recruitment centers

1

Recruitment center
Name of recruitment center
Labafenezhad Hospital
Full name of responsible person
Fatemeh Sadat hosaini
Street address
Valley 9, Pasdaran street, Tehran.
City

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Vice Chancellor for research of Kashan University of Medical Sciences

Full name of responsible person

Dr. Gholamali Hamidi

Street address

Ghotb ravandi bolvard, Kashan

City

Kashan

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 of Kashan 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

"Kashan University of Medical Sciences"

Full name of responsible person

Fatemeh Sadat Hosaini

Position

Bachelor Of Science

Other areas of specialty/work**Street address**

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Person responsible for scientific inquiries

Contact

Name of organization / entity

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Full name of responsible person

Dr. Mohsen Adib-Hajbaghery

Position

Professor, faculty member

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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