

Clinical Trial Protocol

Iranian Registry of Clinical Trials

29 Jun 2026

Comparison the effect of aromatherapy with lavender and rose on anxiety before surgery

Protocol summary

Summary

The purpose of this study is to Comparison the effect of aromatherapy with lavender and rose on anxiety before surgery. Inclusion criteria are do not using of other anxiety reduction methods and no history of eczema and allergy to plants and no disorder in smell. Exclusion criteria are request to withdraw from the study at any stage and the occurrence of any effects indicating an allergic reaction to the essential used oils. In this clinical trial 135 patients will be selected according to the inclusion criteria and randomly assigned nine-point reversal blocks to three groups of placebo, lavender and red rose (45 subjects each group). In inhaled aromatherapy groups, two drops of essential oils of each plant will be placed on a cloth and the patient will be inhaled from a distance of 7-10 cm and kept for 20 minutes. In the placebo group, two drops of water will be placed on a tissue and similar to the previous two groups, the patient will be inhaled. Immediately before and after the intervention, the mean anxiety score will be determined using the Spielberger standard anxiety questionnaire.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2017092136309N1**

Registration date: **2017-10-21, 1396/07/29**

Registration timing: **prospective**

Last update:

Update count: **0**

Registration date

2017-10-21, 1396/07/29

Registrant information

Name

Hossein Jeddi

Name of organization / entity

Gonabad University of Medical Sciences

Country

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Email address

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Recruitment status

Recruitment complete

Funding source

Vice chancellor for research, Gonabad University of Medical Sciences

Expected recruitment start date

2017-10-23, 1396/08/01

Expected recruitment end date

2018-01-21, 1396/11/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Comparison the effect of aromatherapy with lavender and rose on anxiety before surgery

Public title

The effect of aromatherapy on anxiety before surgery

Purpose

Supportive

Inclusion/Exclusion criteria

Inclusion criteria: do not using of other anxiety reduction methods; no history of eczema and allergy to plants; no disorder in smell. Exclusion criteria: request to withdraw from the study at any stage; the occurrence of any effects indicating an allergic reaction to the essential used oils.

Age

No age limit

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **135**

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 Gonabad University of Medical Sciences

Street address

Gonabad University of Medical Sciences, Next to the Asian Road, Gonabad, Khorasan Razavi, Iran, Islamic Republic Of

City

Gonabad

Postal code

Approval date

2016-07-20, 1395/04/30

Ethics committee reference number

GMU.REC.1395.43

Health conditions studied

1

Description of health condition studied

Anxiety before surgery

ICD-10 code

F41.1

ICD-10 code description

Generalized anxiety disorder

Primary outcomes

1

Description

Anxiety

Timepoint

Immediately before and after the intervention

Method of measurement

The Spielberger standard instrument (STAI) is widely used by psychologists, specialists and researchers. The credibility and reliability of the Persian translation of that has been reviewed and approved by faculty members of the Shahid Beheshti University of Tehran.

Secondary outcomes

empty

Intervention groups

1

Description

In inhaled aromatherapy red rose groups, two drops of essential oils of red rose will be placed on a cloth and the patient will be inhaled from a distance of 7-10 cm and kept for 20 minutes. Immediately before and after the intervention, the mean anxiety score will be determined using the Spielberger standard anxiety questionnaire.

Category

Treatment - Other

2

Description

In inhaled aromatherapy lavender groups, two drops of essential oils of lavender will be placed on a cloth and the patient will be inhaled from a distance of 7-10 cm and kept for 20 minutes. Immediately before and after the intervention, the mean anxiety score will be determined using the Spielberger standard anxiety questionnaire.

Category

Treatment - Drugs

3

Description

In the placebo group, two drops of water will be placed on a cloth and the patient will be inhaled from a distance of 7-10 cm and kept for 20 minutes. Immediately before and after the intervention, the mean anxiety score will be determined using the Spielberger standard anxiety questionnaire. .

Category

Treatment - Other

Recruitment centers

1

Recruitment center

Name of recruitment center

The 15 khordad hospital

Full name of responsible person
Street address
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h.jeddi@gmu.ac.ir
Web page address

Sponsors / Funding sources

1

Sponsor

Name of organization / entity
Vice chancellor for research, Gonabad University of Medical Sciences
Full name of responsible person
Dr. Ali Mohammadpour
Street address
Gonabad University of Medical Sciences, Next to the Asian Road, Gonabad, Khorasan Razavi, Iran, Islamic Republic Of.
City
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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
Vice chancellor for research, Gonabad 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
Gonabad University of Medical Sciences
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Position
BSc in nursing
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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