

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

28 May 2026

### comparison of aromatherapy with Citrus aurantium and lavender on sexual satisfaction of lactating women referred to health care centers

#### Protocol summary

##### Study aim

Determine the effect of aromatherapy with Citrus aurantium and lavender on the sexual satisfaction of lactating women

##### Design

This randomized triple-blind clinical trial study was performed on 210 qualified lactating woman referred to health care centers of Kermanshah. Participants were randomly assigned to three groups (70 in each group) including two interventional and one control group with a placebo and each participant is assigned a code.

##### Settings and conduct

Place of study is health care centers of Kermanshah and participants receive three times a day and each time, inhaled 2 to 3 drops of essence (lavender or Citrus aurantium) or placebo solution that are poured on forearm area of the hand for 40 days. Study is triple-blind.

##### Participants/Inclusion and exclusion criteria

Inclusion criteria: Placing in breastfeeding (3-6 months postpartum), lack of psychological problems or systemic disease known to affect sex performance, lack of sexual problems in husband according wife's comment, absence of stressful events in the last 6 months, having a health record in health care centers, married and living with a permanent husband, use a reliable method of contraception, lack of using any medication that has an effect on sexual response, lack of disorder of olfactory or sensitivity to herbal medicines. Exclusion criteria: quitting the study, occurrence of side effects of medication and sensitivity to essential oils during the study, occurrence of major emotional event, occurrence of pregnancy during the study, occurrence of mental health problems or serious illness during the study, divorce or death or illness of the husband during the study.

##### Intervention groups

Intervention group 1: intervention with Lavender aroma  
Intervention group 2: intervention with Citrus aurantium

Intervention group 3: intervention with placebo (control group)

##### Main outcome variables

The main outcome is the mean of total sexual satisfaction after intervention in study groups.

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20160427027633N3**

Registration date: **2017-12-17, 1396/09/26**

Registration timing: **registered\_while\_recruiting**

Last update: **2017-12-17, 1396/09/26**

Update count: **0**

##### Registration date

2017-12-17, 1396/09/26

##### Registrant information

##### Name

Foruzan Sharifipour

##### Name of organization / entity

Kermanshah University of Medical Sciences

##### Country

Iran (Islamic Republic of)

##### Phone

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##### Recruitment status

**Recruitment complete**

##### Funding source

Kermanshah University of Medical Sciences

##### Expected recruitment start date

2016-12-24, 1395/10/04

##### Expected recruitment end date

2018-04-21, 1397/02/01

**Actual recruitment start date**

empty

**Actual recruitment end date**

empty

**Trial completion date**

empty

**Scientific title**

comparison of aromatherapy with Citrus aurantium and lavender on sexual satisfaction of lactating women referred to health care centers

**Public title**

comparison of aromatherapy with Citrus aurantium and lavender on sexual satisfaction of lactating women

**Purpose**

Treatment

**Inclusion/Exclusion criteria**

**Inclusion criteria:**

willingness to participate in the study having health dossier in health care centers literacy sufficient to complete of the questionnaire married and lives with her husband permanently use a reliable method of contraception exposure during lactation (6-3 months postpartum) lack of mental health problems or known systemic disease affecting a person's sexual function lack of drug dependence lack of sexual problems in husband according wife's comment lack of addicted husband the absence of stressful events in the last 6 months (divorce of parents, death of family members, etc.) the absence of diseases, , lack of stress factors (Parent divorce in the past six months), I the lack of any effective drug on a person's sexual response (Antihypertensive drugs, Thiazide Diuretics, Antidepressants, Antihistamines, Barbiturates, Narcotics, Diazepins, Amphetamines, Cocaine) lack of disorder of olfactory or sensitivity to herbal medicines according wife's comment and other disorders related to nose (fracture, deviated septum, rhinitis and sinusitis

**Exclusion criteria:**

dissuasion of participating in the study incidence of side effects and sensitivity to essential oils during the study emotional major event such as a death of One of the relatives in particular loss of child proof of pregnancy occurrence during the study the incidence of mental health problems or serious illness during the study living away from his husband catching the debilitating diseases such as cancer that person will have to leave the study surgery during the intervention divorce, death or illness of husband during the intervening period of study

**Age**

From **18 years** old to **40 years** old

**Gender**

Female

**Phase**

N/A

**Groups that have been masked**

- Participant
- Investigator
- Data analyser

**Sample size**

Target sample size: **210**

**Randomization (investigator's opinion)**

Randomized

**Randomization description**

In this study, Simple randomization method is used for random allocation (hiding allocation). In this way, 210 cards matched in appearance are prepared and 70 of them have the number or code 1 that specifies the group "Intervention based on essence of Citrus aurantium " and on 70 others of them the number or code 2 which specifies the group "Intervention based on essence of lavender " and on the other 70 Of them, the number or code 3, which specifies the control group, means "routine actions" is written. Then, each eligible person for inclusion in the study randomly takes one of these cards with the codes written on it. In this way, the random allocation of patients to each group is determined without letting the participants to know the nature of the numbers 1, 2, or 3 in the type of intervention that will be assigned.

**Blinding (investigator's opinion)**

Triple blinded

**Blinding description**

In this study, to hide the allocation of aromas (essence of Citrus aurantium or lavender) and placebo (Firouz baby oil) Opaque envelopes sequentially numbered are used. Each envelope contains a glass of essence of Citrus aurantium or lavender or placebo. Medications were placed on the envelope in the randomized allocation sequence by the person not involved in the study. Investigating individuals to study and delivering envelopes are done by the researcher himself from 1 to 210. Therefore, in this study, participant and researcher blindness are used. Blindness of data analyzer also runs. Thus, individuals in two groups of intervention (Citrus aurantium or lavender) and the placebo group (Firouz babyoil) are characterized by code (for example, codes 1, 2 and 3) and data analyzer is not aware of the subject code. So the study is triple-blind.

**Placebo**

Used

**Assignment**

Parallel

**Other design features**

**Secondary Ids**

empty

**Ethics committees**

**1**

**Ethics committee**

**Name of ethics committee**

Ethics committee of Kermanshah University of Medical Sciences

**Street address**

Shahid Beheshti Avenue, Research Deputy of Kermanshah University of Medical Sciences

**City**

Kermanshah

**Province**

Kermanshah

**Postal code**

6715847141

**Approval date**

2017-07-19, 1396/04/28

**Ethics committee reference number**

KUMS.REC.1396.189

## Health conditions studied

### 1

**Description of health condition studied**

sexual satisfaction

**ICD-10 code**

**ICD-10 code description**

## Primary outcomes

### 1

**Description**

sexual satisfaction

**Timepoint**

Before and 40 days after the intervention

**Method of measurement**

Lindaberg's Sexual Satisfaction Questionnaire

## Secondary outcomes

### 1

**Description**

Side effects

**Timepoint**

40 days after start the intervention

**Method of measurement**

Researcher made questionnaire

## Intervention groups

### 1

**Description**

Interventions 1: intervention with Lavender aroma: They received three times a day and each time, 2 to 3 drops of solution are injected into the forearm area of the hand for 40 days in an inhaler.

**Category**

Treatment - Drugs

### 2

**Description**

Interventions 2: intervention with Citrus aurantium aroma: They received three times a day and each time, 2 to 3 drops of solution are injected into the forearm area of the hand for 40 days in an inhaler.

**Category**

Treatment - Drugs

### 3

**Description**

Intervention 3: control with Firuze baby oil aroma: They received three times a day and each time, 2 to 3 drops of solution are injected into the forearm area of the hand for 40 days in an inhaler.

**Category**

Treatment - Drugs

## Recruitment centers

### 1

**Recruitment center**

**Name of recruitment center**

Kermanshah City Health Center

**Full name of responsible person**

fruzan sharifipour

**Street address**

Shahid Beheshti Ave

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## Sponsors / Funding sources

### 1

**Sponsor**

**Name of organization / entity**

Kermanshah University of Medical Sciences

**Full name of responsible person**

susan heydarpour

**Street address**

Dolat Abad, Isar Square

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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**

Kermanshah 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**

School of Nursing and Midwifery

**Full name of responsible person**

Foruzan Sharifipour

**Position**

Master of Science

**Latest degree**

Master

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Susan heydarpour

**Position**

Assistant Professor

**Latest degree**

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**Other areas of specialty/work**

Midwifery

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Kermanshah

**Province**

Kermanshah

**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