

Clinical Trial Protocol

Iranian Registry of Clinical Trials

13 Sep 2026

A Comparison Study of the Effect of foot reflexology with and without violet oil on fatigue, pain after cesarean section in patients

Protocol summary

Study aim

Determining the comparison of the effect of plantar reflexology with and without violet oil on fatigue and pain after cesarean section in patients referred to Tohid Jam Hospital in Bushehr, 2021

Design

Clinical trial with control group, with parallel groups, one-way blind, randomized, on 126 patients. Randomizer software was used for randomization.

Settings and conduct

From the patients referred to Tohid Jam hospital in Shiraz, based on the inclusion criteria, 126 samples were selected by convenience sampling method and randomly assigned to a control group and two intervention groups using Randomizer web-based software. One-way blinding and intervention and control groups are not aware of the type of intervention. Sealed consecutive matte envelopes were used to conceal the allocation.

Participants/Inclusion and exclusion criteria

Inclusion criteria: Being 18-45 years old; Literacy; Perform cesarean section with spinal anesthesia; Complete pregnancy and healthy baby and high Apgar score of the first minute; No allergies to used oils; No skin problems in the massage area; Do not use narcotic sedatives in the last eight hours; Do not be under general anesthesia. Exclusion criteria: Having severe postoperative complications such as excessive bleeding, infection, eclampsia, hospitalization in intensive care units; Reluctance to continue cooperation during stages 2 and 3.

Intervention groups

Group 1: In the first group, the reflexology intervention of the sole of the foot will be performed with violet oil. Second group: In the second group, the reflexology intervention of the sole of the foot will be performed without violet oil. The third group: in this group are cesarean section patients who are not subjected to any intervention.

Main outcome variables

Fatigue, pain

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20211204053272N1**

Registration date: **2022-01-01, 1400/10/11**

Registration timing: **prospective**

Last update: **2022-01-01, 1400/10/11**

Update count: **0**

Registration date

2022-01-01, 1400/10/11

Registrant information

Name

Esmat Mardani

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 71 3647 4254

Email address

esmatmardani6609@gmail.com

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2022-01-21, 1400/11/01

Expected recruitment end date

2022-03-21, 1401/01/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty	
Scientific title	A Comparision Study of the Effect of foot reflexology with and without violet oil on fatigue, pain after cesarean section in patients
Public title	The effect of plantar reflexology on fatigue and pain after cesarean section
Purpose	Treatment
Inclusion/Exclusion criteria	Inclusion criteria: Being 18-45 years old Literacy Perform cesarean section with spinal anesthesia Complete pregnancy and healthy baby and high Apgar score of the first minute No allergies to used oils No skin problems in the massage area Do not use narcotic sedatives in the last eight hours Do not be under general anesthesia Exclusion criteria: Having severe postoperative complications such as excessive bleeding, infection, eclampsia, hospitalization in intensive care units Reluctance to cooperation during stages 2 and 3
Age	From 18 years old to 45 years old
Gender	Female
Phase	1-2
Groups that have been masked	Participant
Sample size	Target sample size: 126
Randomization (investigator's opinion)	Randomized
Randomization description	Randomization is done in a simple and individual way using Randomizer random allocation software. Individuals are listed according to the national code and then divided into groups 1 (reflexology with violet oil), group 2 (reflexology without oil) and group 3 (control) using random allocation software.
Blinding (investigator's opinion)	Single blinded
Blinding description	Only the researcher knows which patient is in the control group and which is receiving reflexology with oil or without oil. Patients are not aware of their groups.
Placebo	Not used
Assignment	Parallel
Other design features	
Secondary Ids	empty

Ethics committees	
1	
Ethics committee	
Name of ethics committee	Ethics committee of Shiraz University of Medical Sciences
Street address	Zand St., in front of Palestine St., the central building of Shiraz University of Medical Sciences, Shiraz,Iran
City	Shiraz
Province	Fars
Postal code	۷۱۳۳۶ - ۷۱۳۴۸
Approval date	2021-09-25, 1400/07/03
Ethics committee reference number	IR.SUMS.NUMIMG.REC.1400.036

Health conditions studied	
1	
Description of health condition studied	Cesarean section
ICD-10 code	O75.82
ICD-10 code description	Onset (spontaneous) of labor after 37 completed weeks of gestation but before 39 completed weeks gestation, with delivery by (planned) cesarean section

Primary outcomes	
1	
Description	Fatigue
Timepoint	Before studying and one hour after studying
Method of measurement	Multidimensional Fatigue Inventory scale
2	
Description	Pain
Timepoint	Before studying and one hour after studying
Method of measurement	Visual Analog Scale

Secondary outcomes	empty
Intervention groups	

1

Description

Intervention group 1: The reflexology intervention of the sole of the foot will be performed with violet oil. After washing the patients's soles and drying them while lying down, the researcher will perform a foot reflexology with 3-5 cc of violet oil for 40 minutes (20 minutes for each foot). The number of intervention sessions will be four times a week for two consecutive days (twice a day at intervals of 6-8 hours). The intervention begins after the patient has RBR (8 to 10 hours after cesarean section) and subsequent reflexology sessions will be performed at approximately a fixed time interval. Diclofenac suppositories are given at each stage of the study if severe pain and analgesia are needed.

Category

Treatment - Drugs

2

Description

Intervention group 2: The foot reflexology intervention will be performed without violet oil. After washing the patients's soles and drying them while lying down, the researcher will perform foot reflexology without violet oil for 40 minutes (20 minutes for each foot). The number of intervention sessions will be four times a week for two consecutive days (twice a day at intervals of 6-8 hours). The intervention begins after the patient has RBR (8 to 10 hours after cesarean section) and subsequent reflexology sessions will be performed at approximately a fixed time interval. Diclofenac suppositories are given at each stage of the study if severe pain and analgesia are needed.

Category

Treatment - Other

3

Description

Control group: In this group are cesarean section patients who are not subjected to any intervention. Patients in this group are only monitored. Diclofenac suppositories are given at each stage of the study if severe pain and analgesia are needed.

Category

N/A

Recruitment centers

1

Recruitment center

Name of recruitment center

Tohid Jam Hospital in Bushehr

Full name of responsible person

Esmat Mardani

Street address

Jam city, Tohid town, Tohid Jam hospital, Bushehr, Iran

City

Bushehr

Province

Boushehr

Postal code

7558174946

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Esmatmardani6609@gmail.com

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Shiraz University of Medical Sciences

Full name of responsible person

Dr. Fatemeh Ghavi

Street address

Namazi Square "Namazi Hospital" Fatemeh School of Nursing and Midwifery, Shiraz, Iran

City

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Fars

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ghavi.fatemeh@yahoo.com

Grant name

Grant code / Reference number

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

Yes

Title of funding source

Shiraz University of Medical Sciences

Proportion provided by this source

65

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

Shiraz University of Medical Sciences

Full name of responsible person

Fatemeh Ghavi

Position

Instructor of Midwifery

Latest degree

Ph.D.

Other areas of specialty/work

Midwifery

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Person responsible for scientific inquiries

Contact

Name of organization / entity

Shiraz University of Medical Sciences

Full name of responsible person

Fatemeh Ghavi

Position

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Latest degree

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Other areas of specialty/work

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Person responsible for updating data

Contact

Name of organization / entity

Shiraz University of Medical Sciences

Full name of responsible person

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Position

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Sharing plan

Deidentified Individual Participant Data Set (IPD)

Yes - There is a plan to make this available

Study Protocol

Yes - There is a plan to make this available

Statistical Analysis Plan

Yes - There is a plan to make this available

Informed Consent Form

Yes - There is a plan to make this available

Clinical Study Report

Yes - There is a plan to make this available

Analytic Code

Not applicable

Data Dictionary

Not applicable

Title and more details about the data/document

Article

When the data will become available and for how long

2022

To whom data/document is available

Researchers working in academic and scientific institutions

Under which criteria data/document could be used

1- Reference in articles and dissertations 2- Having a code of ethics in your studies 3- Valid academic emails

From where data/document is obtainable

Esmat Mardani Hazrat Fatemeh Shiraz School of Nursing and Midwifery Esmatmardani6609@gmail.com 00989173894438

What processes are involved for a request to access data/document

Communication via valid email

Comments