

Clinical Trial Protocol

Iranian Registry of Clinical Trials

29 May 2026

Comparison the effect of hot bag water and eflorage massage in active phase of labor, on Pain severity in primiparous women

Protocol summary

Summary

The aim of study: Comparison the effect of hot bag water and eflorage massage in active phase of labor, on Pain severity in primiparous women. A Randomized (The use of sealed envelopes); not blinded, controlled, single-center, phase two trial. Inclusion criteria: the prediction of vaginal delivery with a normal history and physical examination by a medical team. . Exclusion criteria: Any reasons during the Labour, Start out of the labor and requiring special care or having an emergency cesarean section, Sensitivity to oil, Unwillingness to continue participation in the study. Sample size: 90. Type of Intervention: hot bag water, eflorage massage. Time involved: the active phase of labor and transitional. outcomes: reducing labor pain.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2015090615338N3**

Registration date: **2015-10-19, 1394/07/27**

Registration timing: **registered_while_recruiting**

Last update:

Update count: **0**

Registration date

2015-10-19, 1394/07/27

Registrant information

Name

Shahnaz Golian Tehrani

Name of organization / entity

Nursing and Midwifery faculty

Country

Iran (Islamic Republic of)

Phone

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Email address

golian@tums.ac.ir

Recruitment status

Recruitment complete

Funding source

Tehran University of Medical Science

Expected recruitment start date

2015-09-23, 1394/07/01

Expected recruitment end date

2015-12-22, 1394/10/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Comparison the effect of hot bag water and eflorage massage in active phase of labor, on Pain severity in primiparous women

Public title

reducing labor pain

Purpose

Supportive

Inclusion/Exclusion criteria

Inclusion criteria: primiparous women with physical and mental health patients in the active phase; the prediction of vaginal delivery with a normal history and physical examination by a medical team; The Midwifery no problems in accepting the views of abnormal embryos, multiple pregnancies, fetal macrosomia, and pelvic disproportion, polyhydramnios, oligohydramnios, preeclampsia, premature placental separation, placenta previa, fast delivery and uterine infections; with a singleton pregnancy, cephalic view; no risk of hypertension, diabetes and other medical disorders in pregnancy; No risk of skin diseases like eczema, and any wound in the hot bag and massage; Having spontaneous

contractions. Exclusion criteria: All mothers of the natural course of maternal or fetal Labor Start due to factors such as detachment, cord prolapse, fetal distress out or for any reason during the Labor Start out of the labor and require special care or have an emergency cesarean section; Lack of desire to continue participating in the study; Use of drugs such as oxytocin labor catalyst; Use of an analgesic such as pethidine.

Age

From **18 years** old to **35 years** old

Gender

Female

Phase

2

Groups that have been masked

No information

Sample size

Target sample size: **90**

Randomization (investigator's opinion)

Randomized

Randomization description

Blinding (investigator's opinion)

Not blinded

Blinding description

Placebo

Not used

Assignment

Parallel

Other design features

Tehran University of Medical Science

Secondary Ids

1

Registry name

golian

Secondary trial Id

46po

Registration date

1394-07-02, 777/01-1356

Ethics committees

1

Ethics committee

Name of ethics committee

Tehran University of Medical Sciences

Street address

Central apartment of Tehran University of Medical Sciences, Keshavarz blvd

City

Tehran

Postal code

Approval date

2014-06-30, 1393/04/09

Ethics committee reference number

1525/130/d/93

Health conditions studied

1

Description of health condition studied

delivery

ICD-10 code

O00-O99

ICD-10 code description

Other obstetric conditions, not elsewhere classified

Primary outcomes

1

Description

Pain

Timepoint

Every 30 minutes

Method of measurement

Visual analogue of pain

Secondary outcomes

empty

Intervention groups

1

Description

Group 1:hot water bag for 15 minutes

Category

Other

2

Description

Group 2: eflorage massage for 20 minutes

Category

Other

3

Description

Control group: routine delivery room care

Category

N/A

Recruitment centers

1

Recruitment center

Name of recruitment center

Comprehensive women's hospital

Full name of responsible person

Zahra Kohzad

Street address

Karimkhan Zand Street, Nejatollahi

Street,Comprehensive women's hospital

City

Tehran

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Tehran University of Medical Science research

Full name of responsible person

Dr. Masood Younesian

Street address

Central Building of Tehran University of Medical Sciences, the Qods Street, Keshavarz Blvd

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Tehran

Grant name**Grant code / Reference number****Is the source of funding the same sponsor organization/entity?**

Yes

Title of funding source

Tehran University of Medical Science research

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

Tehran University of Medical Science

Full name of responsible person

Golian tehrani Shahnaz

Position

Faculty member, Master of Midwifery

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Web page address

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