

Clinical Trial Protocol

Iranian Registry of Clinical Trials

09 Jun 2026

Evaluation of the effect of saffron oral capsules on pain severity and duration of the active phase of labor first stage

Protocol summary

Summary

Triple blind study to determine the effect of oral capsules of saffron on the severity and duration of the active phase of labor. The main inclusion criteria: nulliparous , 18-35 years, body mass index 5/19 to 30, gestational age 37-42 weeks, dilatation 3-4 cm cervical, cephalic presentation. Main exclusion criteria: using any herbal drug during the past 48 hours, addiction or using a variety of tobacco and alcohol, the risk of pregnancy complications, or history of any systemic illness or mental disease, the fetus with maternal pelvic disproportion, abnormal fetal heart rate, obstetric complications , cesarean indication, allergic reaction to the saffron Study Population, all nulliparous women that go to the hospital for delivery. Sample size:60. In the beginning of the study, one capsule (Saffron or placebo which is determined with a code), is given to the subjects for oral consumption. Pain severity was measured each hour until the end of active phase of first stage. Uterine contraction pattern is recorded every 30 minutes using tocodinamometer. Considering to the pattern of uterine contractions, fetal heart rate and the mother's vital signs, drug dose was repeated every 2 hours up to a maximum of three doses. If the dilation was according to the pattern of uterine contractions, the next dose of capsule is not used. If uterine contractions were hypertone , poor progress in labor or need to pain relief by pharmacological methods, appropriate measurement is performed and she is excluded from study

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT201205229830N1**

Registration date: **2014-03-10, 1392/12/19**

Registration timing: **retrospective**

Last update:

Update count: **0**

Registration date

2014-03-10, 1392/12/19

Registrant information

Name

Sedigheh Ahmadi

Name of organization / entity

Mashhad University Of Medical Sciences

Country

Iran (Islamic Republic of)

Phone

+98 51 1859 1511

Email address

ahmadis901@mums.ac.ir

Recruitment status

Recruitment complete

Funding source

Vice-chancellor of Mashhad University of Medical Sciences

Expected recruitment start date

2013-11-06, 1392/08/15

Expected recruitment end date

2014-03-06, 1392/12/15

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Evaluation of the effect of saffron oral capsules on pain severity and duration of the active phase of labor first stage

Public title

Evaluation of the effect of saffron oral capsules on pain severity and labor duration

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria: Completing the written informed consent to participate in the study; age of 18-35 years; body mass index of 19.5- 30; gestational age of 37-42 weeks; having no history of pregnancy more than 20 weeks of gestation; singleton pregnancy; cervical dilatation of 3-4 cm; cephalic presentation. Exclusion criteria: using any herbal drug during the past 48 hours; addiction or using a variety of tobacco and alcohol; maternal pregnancy complications (preeclampsia, bleeding during pregnancy, threat to abortion, premature rupture of fetus membranes more than 12 hours before admission of mother), having or history of any systemic disease (diabetes, hypertension, heart and kidney disease, etc), history of mental disorder, speaking, hearing and mental disorder, disproportion of maternal pelvic with fetal head; estimated fetal weight less than 2500 g and more than 4000 g (according to Johnson rules); fetal heart rate abnormalities; any abnormalities in the fetus (based on sonography; obstetric problems; multiple pregnancy; previous surgery on the uterus; cervix and delivery canal; cesarean indication; allergic reaction to the saffron

Age

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

Gender

Female

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **60**

Randomization (investigator's opinion)

Not randomized

Randomization description

Blinding (investigator's opinion)

Triple blinded

Blinding description

Placebo

Used

Assignment

Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Mashhad University Of Medical Sciences

Street address

Mashhad University Of Medical Sciences

City

Mashhad

Postal code

8564917794

Approval date

2013-06-22, 1392/04/01

Ethics committee reference number

911184

Health conditions studied

1

Description of health condition studied

delivery

ICD-10 code

O80.8

ICD-10 code description

Other single spontaneous delivery

Primary outcomes

1

Description

pain intensity of labor

Timepoint

1 hour intervals during the intervention.

Method of measurement

Pain intensity using a visual pain scale

2

Description

Duration of the active phase of labor first stage

Timepoint

1-2 hour intervals during the intervention

Method of measurement

During the active phase of labor and vaginal exam registration

Secondary outcomes

1

Description

Uterine contractions

Timepoint

During each 30-minute intervention

Method of measurement

Tocodynamomtr device and touch the hand

Intervention groups

1

Description

Capsules 250 mg of the drug (saffron) every 2 hours and up to 3 doses, the intensity of uterine contractions and the progress of labor is consumed orally by the research unit

Category

Placebo

2

Description

Capsules 250 mg of the placebo every 2 hours and up to 3 doses, the intensity of uterine contractions and the progress of labor is consumed orally by the research unit

Category

Placebo

Recruitment centers

1

Recruitment center

Name of recruitment center

Seventeen Hospitals September Mashhad - Social Security Organization

Full name of responsible person

Azhari - MSc in Nursing - Faculty of Nursing and Midwifery, Mashhad

Street address

Mashhad University Of Medical Sciences

City

Mashhad

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Research Council of Mashhad University of Medical Sciences

Full name of responsible person

sedigheh Azhari

Street address

Mashhad University Of Medical Sciences

City

Mashhad

Grant name

Grant code / Reference number

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

Yes

Title of funding source

Research Council of Mashhad 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

Mashhad University of Medical Sciences

Full name of responsible person

Sedigheh Ahmadi

Position

Student Master of Midwifery

Other areas of specialty/work

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Contact

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Full name of responsible person

Sedigheh Ahmadi

Position

Graduate student Midwifery

Other areas of specialty/work

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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