

Clinical Trial Protocol

Iranian Registry of Clinical Trials

27 Jul 2026

The effect of low dose Bupivacaine with fentanyl in spinal anesthesia for cesarean delivery to prevent maternal hypotension

Protocol summary

Summary

The aim of this randomized double blind clinical trial is reducing usual dose of bupivacaine for spinal anesthesia by adding fentanyl. Can be reducing common side effects of bupivacaine (hypotension), this result in an appropriate level of anesthesia in patients. 105 pregnant women (18 to 40-year-old) candidate for elective cesarean section with spinal anesthesia entered the study. Who had singleton term pregnancies in class I physical health classification of the American Society of Anesthesia (ASA). And mothers with a history of hypertension, diabetes, cardiovascular and cerebral diseases, known fetal anomaly, any kind of contraindications for spinal anesthesia and those with difficult tracheal intubation were excluded. They were randomly divided into three groups of 35 women. The first group received 10 mg of bupivacaine 0.5%, the second received 8 mg of bupivacaine 0.5% with 25 microgram of fentanyl, and the third received 8 mg of bupivacaine 0.5% with 0.4 ml of normal saline Intrathecally. Blood pressure, heart rate, sensory level, degree of motor block, pain score during operation, nausea and vomiting, time taken to recover from sensory and motor block were measured and compared in three groups during and after surgery. Our data will be analyzed using the using t-test, Mann-Whitney test and Chi-square tests. A P value <0.05 will be considered statistically significant.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT201704183305N8**

Registration date: **2017-04-28, 1396/02/08**

Registration timing: **retrospective**

Last update:

Update count: **0**

Registration date

2017-04-28, 1396/02/08

Registrant information

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Name of organization / entity

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Recruitment status

Recruitment complete

Funding source

Vice-chancellor Of Research Babol University Of Medical Sciences

Expected recruitment start date

2015-12-06, 1394/09/15

Expected recruitment end date

2016-12-05, 1395/09/15

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The effect of low dose Bupivacaine with fentanyl in spinal anesthesia for cesarean delivery to prevent maternal hypotension

Public title

The Effect Of Spinal Anesthesia Drug to Prevent Maternal Hypotention

Purpose

Prevention

Inclusion/Exclusion criteria
Inclusion criteria: pregnant women, candidate for elective CS, in class I physical health classification of the American Society of Anesthesia (ASA). All women had singleton term pregnancies. Exclusion criteria: Mothers with a history of hypertension, diabetes, cardiovascular and cerebral diseases, known fetal anomaly, any kind of contraindications for spinal anesthesia, patients weighing over 100 kg and those with difficult tracheal intubation.

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

Gender
Female

Phase
N/A

Groups that have been masked
No information

Sample size
Target sample size: **105**

Randomization (investigator's opinion)
Randomized

Randomization description

Blinding (investigator's opinion)
Double blinded

Blinding description

Placebo
Not used

Assignment
Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethics committee of Babol University Of Medical Sciences

Street address

Babol University of Medical Sciences, Ganjafrooz Street, Babol

City

Babol

Postal code

47176-41367

Approval date

2015-12-06, 1394/09/15

Ethics committee reference number

MUBABOL.REC.1394.232

Health conditions studied

1

Description of health condition studied

Maternal hypotension

ICD-10 code
O26.5

ICD-10 code description
maternal hypotension syndrome

Primary outcomes

1

Description

Maternal blood pressure

Timepoint

Maternal blood pressure was monitored and recorded every two minutes for the first 15 minutes following spinal anesthesia, and then every 5 minutes until completion of surgery.

Method of measurement

Non invasive blood pressure cuff

Secondary outcomes

1

Description

Sensory level and motor block

Timepoint

After spinal anesthesia

Method of measurement

Sensory level was assessed using the tip of a needle sterilized with alcohol cotton, and motor block was assessed and recorded using Bromage Scale (BS) (BS=0, full knee and foot flexion; BS=1, ability to move knee; BS=2, ability to move feet only; BS=3, full motor block).

2

Description

pain

Timepoint

Pain level during surgery

Method of measurement

VAS (Visual analogues Scale)

3

Description

Return of sensory and motor block

Timepoint

After Surgery

Method of measurement

Examination, Question & Observation

Intervention groups

1

Description

Control group: 10 mg 0.5% bupivacaine was injected into subarachnoid space.

Category

Prevention

2

Description

Intervention Group1: 8 mg of bupivacaine 0.5 with 25 micrograms of fentanyl was injected into subarachnoid space.

Category

Prevention

3

Description

Intervention Group2: 8 mg of bupivacaine 0.5% with 0.4 micrograms of Normal saline was injected into subarachnoid space

Category

Prevention

Recruitment centers

1

Recruitment center

Name of recruitment center

Rouhani Hospital of Babol

Full name of responsible person

Dr Hakimeh Alereza Amiri

Street address

Rouhani Hospital of Babol, Ganjafroz St.

City

Babol

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Vice chancellor research for Babol University of Medical Sciences

Full name of responsible person

Dr. Ali Akbar Moghadamnia

Street address

Babol University Of Medical Sciences, Ganjafrooz street, Babol

City

Babol

Grant name

Grant code / Reference number

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

Yes

Title of funding source

Vice chancellor research for Babol 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

Babol University of Medical Sciences

Full name of responsible person

Dr. Hakimeh Alereza Amiri

Position

Assistant Professor of Anesthesiology

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