

Clinical Trial Protocol

Iranian Registry of Clinical Trials

19 Jul 2026

Evaluation of intrathecal dexmedetomidine effect on block characteristics during spinal anesthesia for cesarean section

Protocol summary

Summary

Objective: Evaluation of effect of intrathecal co-administration of dexmedetomidine and bupivacaine on spinal anesthesia blocks characteristics in patient undergoing elective caesarean section . Setting : In this prospective randomized double blind clinical trail 50 patients who will scheduled for cesarean section under spinal anesthesia in besat hospital of sanandaj will be randomly divided in to 2 equal 25 patient group . Inclusion criteria: writing consent form, American Society of Anesthesiologist physical status 1 or 2. Exclusion criteria: preeclampsia or eclampsia, diabetes mellitus, chronic hypertension, cardiovascular disease, renal disease, hepatic failure, and known allergy to study drugs. Method: After insertion of two G20 venous catheters in dorsum of patient's hands 10 ml/Kg normal saline will be infused to patients. Standard monitoring will be start for patient and basic values will be record. Spinal anesthesia will be done in sitting position via L3-L4 or L4-L5 space using disposable (G=26) needle. Following spinal anesthesia patient immediately turned to supine position and about 5 degree head down. Intervention: induction of spinal anesthesia will be done by using 12.5 mg bupivacaine plus 5 mic dexmedetomidine in study group, and the same value of bupivacaine plus 0.5 ml 0.9% normal saline in control group (volume of drug will be reached to 3 ml by adding stilled water in both group). Primary outcome: peak of sensory block and the needing time to reach that, the needing time to reach motor block to highest degree of modified bromage scale, time of regression of sensory block to level of S1 and regression of motor block to modified bromage scale 1. Secondary outcome: cardiovascular change during first 30 min of anesthesia, and complication of block. Blinding: patient, anesthesiologist who do spinal anesthesia, and the anesthesiologist who evaluate study outcomes are unaware about patients grouping .

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT201512051766N8**

Registration date: **2016-01-20, 1394/10/30**

Registration timing: **registered_while_recruiting**

Last update:

Update count: **0**

Registration date

2016-01-20, 1394/10/30

Registrant information

Name

Karim Nasseri

Name of organization / entity

Kurdistan University of Medical Sciences

Country

Iran (Islamic Republic of)

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Email address

nasseri_k@muk.ac.ir

Recruitment status

Recruitment complete

Funding source

Kurdistan University of Medical Sciences .Vice chancellor for research

Expected recruitment start date

2015-05-22, 1394/03/01

Expected recruitment end date

2016-06-21, 1395/04/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Evaluation of intrathecal dexmedetomidine effect on block characteristics during spinal anesthesia for cesarean section

Public title

Evaluation of intrathecal dexmedetomidine effect on block characteristics during spinal anesthesia for cesarean section

Purpose

Prevention

Inclusion/Exclusion criteria

Inclusion criteria : American Society of Anesthesiologist Physical status 1 or 2 Exclusion criteria : Any absolute or relative contraindication for spinal anesthesia ; BMI > 25 ; history of taking α_2 antagonist drugs ; history of hypertension or antihypertensive drug taking ; history of heart disease (ischemic ,valular ,blocks) ; renal failure ; hepatic disease ; history of allergy to study drugs ; inadequate spinal block level

Age

From **18 years** old to **45 years** old

Gender

Female

Phase

2-3

Groups that have been masked

No information

Sample size

Target sample size: **50**

Randomization (investigator's opinion)

Randomized

Randomization description**Blinding (investigator's opinion)**

Double blinded

Blinding description**Placebo**

Used

Assignment

Parallel

Other design features

.

Secondary Ids

empty

Ethics committees**1****Ethics committee****Name of ethics committee**

Kurdistan University of Medical Sciences ethic committee

Street address

Pardis of medicien faculty , Pasdaran BLV

City

Sanandaj

Postal code**Approval date**

2015-08-15, 1394/05/24

Ethics committee reference number

MUK.REC.1394.141

Health conditions studied**1****Description of health condition studied**

spinal anesthesia

ICD-10 code

Other comp

ICD-10 code description

O29.8

Primary outcomes**1****Description**

peak of sensory block

Timepoint

every 15 min until reaching equal result inthree consecutive session

Method of measurement

using Cotton alcohol method in midclavicular line from distal to proximal of body

2**Description**

needing time to reach the peak of sensory block

Timepoint

every 2 min until reaching equal result in three consecutive session

Method of measurement

using Cotton alcohol method in midclavicular line from distal to proximal of body

3**Description**

needing time to reach motor block to highest degree

Timepoint

every 15 min

Method of measurement

modified bromage scale

4**Description**

time of regression of sensory block

Timepoint

every 15 min in post anesthesia care unit

Method of measurement

time need to regression of sensory block to level of S1 using

5**Description**

time need to regression of motor block

Timepoint

every 15 min in post anesthesia care unit
Method of measurement
based on modified bromage scale

Secondary outcomes

1

Description
cardiovascular chang
Timepoint
every 10 min
Method of measurement
Using indirect blood pressure measurement via
automated devise

Intervention groups

1

Description
Intervention : 12.5 Mg bupivacaine 0.5% plus 5 mic
dexmedetomidine (total volume 3 MI by adding distilled
water)
Category
Treatment - Drugs

2

Description
Control : 12.5 Mg bupivacaine 0.5% plus 0.5 MI 0.9%
normal saline (total volume 3 MI)
Category
Placebo

Recruitment centers

1

Recruitment center
Name of recruitment center
operating room number2, Besat Hospital
Full name of responsible person
Dr.KarimNasseri
Street address
Besat hospital , Keshavarz Street
City
Sanandaj

Sponsors / Funding sources

1

Sponsor
Name of organization / entity
Kurdistan University of Medical Sciences .Vice
chancellor for research
Full name of responsible person
Ebrahim Ghaderi
Street address
Pardis of University

City
Sanandaj
Grant name
Grant code / Reference number
**Is the source of funding the same sponsor
organization/entity?**
Yes
Title of funding source
Kurdistan University of Medical Sciences .Vice chancellor
for 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
Sanandaj Faculty of Medicine
Full name of responsible person
Dr.Ebrahim Ghaderi
Position
Epidemiologist / Head of research concil of KUMS
Other areas of specialty/work
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Person responsible for scientific inquiries

Contact
Name of organization / entity
Department of Anesthesiokogy , Kurdistan University
of Medicak Sciences
Full name of responsible person
Dr. Karim Nasseri
Position
Associate Professor / Anesthesiologist
Other areas of specialty/work
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Person responsible for updating data

Contact

Name of organization / entity

Sanandaj Faculty of Medicine

Full name of responsible person

Dr. Karim Nasseri

Position

Anesthesiologist / Department of Anesthesiology

Other areas of specialty/work

Street address

Besat Hospital , Keshavarz Street

City

Sanandaj

Postal code

Phone

00

Fax

Email

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