

Clinical Trial Protocol

Iranian Registry of Clinical Trials

11 Sep 2026

Comparison of hypobaric unilateral spinal anesthesia versus general anesthesia on heart rate, blood pressure change and mortality in hip fracture surgery

Protocol summary

Study aim

To determine the comparative effect of unilateral spinal anesthesia versus general anesthesia on hemodynamics and mortality in hip fracture surgery in the elderly.

Design

Controlled clinical trial, with parallel groups, phase 3 on 60 patients.

Settings and conduct

This study will be conducted as a clinical trial on patients over 60 years of age with hip fracture referred to Shahid Rajaei Hospital in Qazvin. In the intervention group, patients will undergo unilateral spinal anesthesia with hypobaric solution injection including ropivacaine and sufentanil in the subarachnoid space. In the control group, patients will undergo general anesthesia with standard drugs including propofol, midazolam, sufentanil, atracurium, and isoflurane. In both groups, mean arterial blood pressure (MAP) and heart rate will be recorded every three minutes before, during, and after the operation.

Participants/Inclusion and exclusion criteria

Inclusion criteria: People over 60 years of age referring to Shahid Rajaei Hospital in Qazvin who are undergoing orthopedic surgery due to a hip fracture. Exclusion criteria: multiple trauma (multiple injuries or damage to other organs), increased intracranial pressure (high ICP), emergency surgery or urgent need for anesthesia, spinal injection site infection, coagulation disorders or high INR, history of stroke (CVA) or myocardial infarction (MI) within the last three months, patients in shock.

Intervention groups

Intervention group: Anesthesia is performed by intra-subarachnoid injection of hypobaric solution containing ropivacaine and sufentanil. Control group: Patients are placed under general anesthesia with standard drugs including propofol, midazolam, sufentanil, atracurium and isoflurane.

Main outcome variables

Incidence of severe intraoperative hypotension, intraoperative norepinephrine and ephedrine intake, seven-day postoperative mortality

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20250809066791N1**

Registration date: **2025-11-04, 1404/08/13**

Registration timing: **prospective**

Last update: **2025-11-04, 1404/08/13**

Update count: **0**

Registration date

2025-11-04, 1404/08/13

Registrant information

Name

Tarokh Rajabinezhad

Name of organization / entity

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2025-11-22, 1404/09/01

Expected recruitment end date

2026-04-21, 1405/02/01

Actual recruitment start date

empty
Actual recruitment end date
empty
Trial completion date
empty
Scientific title
Comparison of hypobaric unilateral spinal anesthesia versus general anesthesia on heart rate, blood pressure change and mortality in hip fracture surgery
Public title
Comparison of hypobaric unilateral spinal anesthesia versus general anesthesia on heart rate, blood pressure change and mortality in hip fracture surgery
Purpose
Treatment
Inclusion/Exclusion criteria
Inclusion criteria:
People over 60 years of age who visit Shahid Rajaei Hospital in Qazvin and undergo orthopedic surgery due to a hip fracture
Exclusion criteria:
Allergy to ropivacaine Multiple trauma
Age
From 60 years old
Gender
Both
Phase
3
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)
Not 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 Qazvin University of Medical Sciences
Street address
Research and Technology deputy ,Mavaddat Alley,Shahid Beheshti Blvd
City

Qazvin
Province
Qazvin
Postal code
3413996134
Approval date
2025-11-02, 1404/08/11
Ethics committee reference number
IR.QUMS.REC.1404.276

Health conditions studied

1

Description of health condition studied

Hip fracture surgery

ICD-10 code

S72.0

ICD-10 code description

Fracture of head and neck of femur

Primary outcomes

1

Description

Severe hypotension during surgery

Timepoint

From the start of anesthesia to the end of the surgery

Method of measurement

Non-invasive blood pressure measurement with standard monitoring every three minutes

2

Description

Intraoperative norepinephrine intake

Timepoint

From the start of anesthesia or numbing to the end of the surgery

Method of measurement

Total dose recorded on the anesthesia sheet

3

Description

Amount of ephedrine taken during surgery

Timepoint

From the start of anesthesia or numbing to the end of the surgery

Method of measurement

Total dose recorded on the anesthesia sheet

4

Description

Seven-day postoperative mortality

Timepoint

From the end of surgery to seven days later

Method of measurement

Clinical follow-up and hospital record review

Secondary outcomes

empty

Intervention groups

1

Description

Intervention group: Unilateral Spinal Anesthesia with Ropivacaine and Sufentanil: In this group, patients undergo Unilateral Spinal Anesthesia (ULSA). After injecting 500 cc of normal saline, the patient is placed in a sitting position. The hypobaric anesthetic solution is prepared with the following ingredients: 0.5% Ropivacaine, 2 ml (10 mg) of Sufentanil, 1 ml (5 µg) of distilled water, 1 ml. A total of 4 cc of the prepared solution is injected into the subarachnoid space L4-L5 (or a higher or lower space) with a 23 G needle. After injection, the patient is placed in a lateral position on the non-involved side (fracture area upwards) for 5 minutes to limit the spread of the drug to that side. In case of mild pain during the procedure (NRS <4), an opioid infusion is used. If the patient's pain is not controlled, general anesthesia is performed and the patient is excluded from the study. Monitoring of mean arterial blood pressure (MAP) and heart rate is performed every 3 minutes, as in the control group, and in case of hypotension, treatment with ephedrine and, if necessary, norepinephrine is performed.

Category

Treatment - Drugs

2

Description

Control group: General anesthesia with isoflurane and standard induction drugs: In this group, patients are put under general anesthesia. After injecting 500 cc of normal saline and premedication with midazolam at a dose of 0.02 mg/kg, sufentanil at a dose of 0.2 µg/kg, propofol at a dose of 1 mg/kg, and atracurium at a dose of 0.5 mg/kg, anesthesia is used to induce anesthesia. After ensuring proper airway conditions and performing endotracheal intubation, the patient's anesthesia is maintained with isoflurane and oxygen/medical air based on the patient's hemodynamic status. During the procedure, mean arterial blood pressure (MAP) and heart rate are monitored every 3 minutes. In case of hypotension, a bolus of ephedrine 5 to 10 mg is first injected and repeated if necessary up to a maximum of 20 mg. If hypotension persists, norepinephrine infusion is initiated at a dose of 5-10 µg/min. At the end of the procedure, the patient is monitored and data are recorded for hemodynamic parameters during recovery.

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Shahid Rajaei hospital

Full name of responsible person

Tarokh Rajabinezhad

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

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Grant name

Grant code / Reference number

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

Yes

Title of funding source

Qazvin University of Medical Sciences

Proportion provided by this source

100

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

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Full name of responsible person
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Sharing plan

Deidentified Individual Participant Data Set (IPD)

Undecided - It is not yet known if there will be a plan to make this available

Study Protocol

Undecided - It is not yet known if there will be a plan to make this available

Statistical Analysis Plan

Not applicable

Informed Consent Form

Undecided - It is not yet known if there will be a plan to make this available

Clinical Study Report

Undecided - It is not yet known if there will be a plan to make this available

Analytic Code

Undecided - It is not yet known if there will be a plan to make this available

Data Dictionary

Undecided - It is not yet known if there will be a plan to make this available