

Clinical Trial Protocol

Iranian Registry of Clinical Trials

07 Jul 2026

A survey on analgesic effects of ultrasonographic transabdominal plane block for postoperative pain control of abdominal hysterectomy

Protocol summary

Summary

Pain and discomfort after abdominal hysterectomy is high and most of it is due to abdominal incision. Aberrant sensory nerves pass from the fascia between abdominal muscles and with trans-abdominal block we can control pain. We usually use epidural or intravenous narcotics to control pain but these ways have side effects like vomiting. With trans-abdominal block we have a simple way to control pain without entering the epidural space. But unfortunately this method is not well recognized by anesthesiologists, so we decided to do this research and declare its results to anesthesiologists. All patients that have gone under abdominal hysterectomy with pfannenstiel incision and don't have systemic and neurologic diseases will be invited to participate in this study. We will put them randomly in two groups. For the first group we will use trans-abdominal block with pain pump and for other group only pain pump will be used. At 2, 6, 12, 24 and 48 hours after operation we will measure score of pain and satisfaction with Visual Analogue Scale test and Global Clinical Improvement and then we will compare the amount of pain scores and the amount of fentanyl is used in the two groups.

General information

Acronym

A survey on analgesic effects of trans abdominal block

IRCT registration information

IRCT registration number: **IRCT2013021012151N1**

Registration date: **2013-03-10, 1391/12/20**

Registration timing: **registered_while_recruiting**

Last update:

Update count: **0**

Registration date

2013-03-10, 1391/12/20

Registrant information

Name

Fariba Almassinokiani

Name of organization / entity

Iran University of Medical Sciences

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Tehran University of Medical Sciences

Expected recruitment start date

2013-02-28, 1391/12/10

Expected recruitment end date

2014-03-01, 1392/12/10

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

A survey on analgesic effects of ultrasonographic transabdominal plane block for postoperative pain control of abdominal hysterectomy

Public title

Effects of transabdominal plane block for postoperative pain control of hysterectomy

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria: All of elective abdominal hysterectomies with pfannenstiel incision will enter to our study. Exclusion criteria: If the patient doesn't accept;

local anesthetic drugs allergy; infection on injection site; coagulation disorders; medical disorders that interfere with interpretation of pain; addiction to illicit drugs or alcohol; liver or renal insufficiency; sever obesity.

Age

From **15 years** old to **45 years** old

Gender

Female

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **64**

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

Tehran University of Medical Sciences

Street address

Poursina street , Enghelab AVE, Tehran , Iran

City

Tehran

Postal code

Approval date

2012-10-08, 1391/07/17

Ethics committee reference number

17543-30-01-91

Health conditions studied

1

Description of health condition studied

Analgesic effect of trans abdominal block on post abdominal hysterectomy pain

ICD-10 code

Z41.8

ICD-10 code description

Other procedures for purposes other than remedying health state

Primary outcomes

1

Description

Severity of post operation pain

Timepoint

2 and 6 and 12 and 24 and 48 hours after operation

Method of measurement

Visual Analogue Scale test

Secondary outcomes

1

Description

Amount of patient's satisfaction from after operation analgesia

Timepoint

2 and 6 and 12 and 24 and 48 hours after operation

Method of measurement

Global Clinical Improvement (GCI) Test

2

Description

The amount of Fantanyle that will be used for pain pump

Timepoint

2 and 6 and 12 and 24 and 48 hours after operation

Method of measurement

Micro geram of fantanyle that will be necessary for pain pump

Intervention groups

1

Description

Intervention group: we use trans abdominal block with pain pump for post abdominal hysterectomy pain control

Category

Other

2

Description

control group : only using pain pump for control of after operation pain

Category

Other

Recruitment centers

1

Recruitment center

Name of recruitment center

Rasoul Akram Hospital

Full name of responsible person

Dr Farnad Imani

Street address

pain ward ; 4th stage ; Rasoul Akram Hospital ;
Niayesh Street ; Sattarkhan Street

City

Tehran

f-almasi@sina.tums.ac.ir

Web page address

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Tehran University of Medical Sciences

Full name of responsible person

Dr Fariba Almassinokiani

Street address

Rasoul Akram Hospital , Niayesh street, Sattarkhan street

City

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 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

Tehran University of Medical Sciences

Full name of responsible person

Dr Fariba Almassinokiani ; Dr Farnad Imani

Position

Obstetrician and Gynecologist ; Anesthesiologist (pain)

Other areas of specialty/work

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Person responsible for scientific inquiries

Contact

Name of organization / entity

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Full name of responsible person

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Position

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Other areas of specialty/work

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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