

Clinical Trial Protocol

Iranian Registry of Clinical Trials

28 Jul 2026

Evaluation of the effects of gabapentin on pain relief after total abdominal hysterectomy

Protocol summary

Summary

Objective: Evaluation of the effects of gabapentin on pain relief after total abdominal hysterectomy. Design: double blind randomized clinical trial. Setting and conduct: candidates for total abdominal hysterectomy will be randomly assigned in two groups of intervention or placebo. In the intervention group; 800 milligrams of oral gabapentin and in the placebo group, placebo will be administered 1 hour before surgery. The method of anesthesia will be the same for both groups. After finishing the surgery and at the recovery room, 30 milligrams of pethidine will be injected for all subjects (point zero) and pain will be evaluated using Visual Analogue Scale (VAS). Then, in 2, 6, 12 and 24 hours after surgery, pain score according to VAS, and the severity of nausea and vomiting will be recorded. Participants: Inclusion criteria: candidates for total abdominal hysterectomy; age between 35- 60 years old. Exclusion criteria: history of known psychiatric disorders; drug abuse; patients with chronic pain. Interventions: In the intervention group; 800 milligrams of oral gabapentin and in the placebo group, placebo will be administered 1 hour before surgery. Main outcome: pain relief.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT201603172624N20**

Registration date: **2016-04-08, 1395/01/20**

Registration timing: **registered_while_recruiting**

Last update:

Update count: **0**

Registration date

2016-04-08, 1395/01/20

Registrant information

Name

Maryam Kashanian

Name of organization / entity

Iran University of Medical Sciences

Country

Iran (Islamic Republic of)

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Email address

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Recruitment status

Recruitment complete

Funding source

Investigator.

Expected recruitment start date

2015-04-07, 1394/01/18

Expected recruitment end date

2016-04-20, 1395/02/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Evaluation of the effects of gabapentin on pain relief after total abdominal hysterectomy

Public title

Pain relief effects of gabapentin after surgery

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria: candidates for total abdominal hysterectomy; age between 35- 60 years old. Exclusion criteria: history of known psychiatric disorders; drug abuse; patients with chronic pain.

Age

From **35 years** old to **60 years** old

Gender

Female

Phase

3

Groups that have been masked

No information

Sample size

Target sample size: 76

Randomization (investigator's opinion)

Randomized

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

Double blinded

Blinding description**Placebo**

Used

Assignment

Parallel

Other design features

Randomization was performed as stratified block randomization.

Secondary Ids

empty

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

Ethics committee of Iran.University of Medical Sciences

Street address

Hemmat highway, Chamran Cross

City

Tehran

Postal code**Approval date**

2014-10-21, 1393/07/29

Ethics committee reference number

93/3393 /105/3

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

Pain after hysterectomy

ICD-10 code

N00-N99

ICD-10 code description

Diseases of the genitourinary system

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

Pain .

Timepoint

2, 6, 12 , 24 hours.

Method of measurement

VAS

Secondary outcomes

empty

Intervention groups**1****Description**

Intervention group: gabapentin administration 1 hour before surgery.

Category

Treatment - Drugs

2**Description**

Control group: placebo administration 1 hour before surgery.

Category

Treatment - Drugs

Recruitment centers**1****Recruitment center****Name of recruitment center**

Akbarabadi Teaching Hospital

Full name of responsible person

Maryam Kashanian

Street address

Molavi avenue, Molavi Cross Road.

City

Tehran

Sponsors / Funding sources**1****Sponsor****Name of organization / entity**

Vice chancellor for research ,Iran University of Medical Sciences.

Full name of responsible person

Dr Seyed Ali Javad Mousavi

Street address

Hemmat High way, Junction of Chamran Highway.

City

Tehran

Grant name**Grant code / Reference number****Is the source of funding the same sponsor organization/entity?**

Yes

Title of funding source

Vice chancellor for research ,Iran 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

Iran University of Medical Sciences

Full name of responsible person

Maryam Kashanian

Position

Professor.

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