

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

28 May 2026

### The post operative analgesic effect of gabapentine in patients undergoing abdominal hysterectomy.

#### Protocol summary

##### Summary

This study is designed to survey gabapentin analgesic effect on post-abdominal hysterectomy pain. Fifty patients aged 35-50 years candidate for abdominal hysterectomy will be included in this study. After signing the informed consent, the patients will be randomly assigned into two groups. The intervention group will receive 1200mg gabapentin and the control group 1200mg placebo. Then, all patients will undergo the operation with the same procedure under the same anesthesia method. The pain will be measured at 2, 6, 12, and 24 hours after the operation with visual analogue scale. Then, the patients will receive 5mg morphine if the pain score was more than 4. Pain intensity, the amount of morphine consumed, nausea, vomiting, and light headedness will be measured and compared between groups.

#### General information

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT138903112967N2**

Registration date: **2010-06-01, 1389/03/11**

Registration timing: **registered\_while\_recruiting**

Last update:

Update count: **0**

##### Registration date

2010-06-01, 1389/03/11

##### Registrant information

##### Name

Fatemeh Foroozfar

##### Name of organization / entity

Kashan University of Medical Sciences

##### Country

Iran (Islamic Republic of)

##### Phone

+98 36 1446 0180

##### Email address

forozaanfar\_f@kaums.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

Vice-chancellor for Research Kashan University of Medical Sciences

##### Expected recruitment start date

2010-05-16, 1389/02/26

##### Expected recruitment end date

2011-03-20, 1389/12/29

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

##### Scientific title

The post operative analgesic effect of gabapentine in patients undergoing abdominal hysterectomy.

##### Public title

The post operative analgesic effect of gabapentine in patients undergoing abdominal hysterectomy.

##### Purpose

Treatment

##### Inclusion/Exclusion criteria

Inclusion criteria: age 35-50 years, candidate for hysterectomy Exclusion criteria: Allergy to morphine or gabapentine, chronic use of analgesic drugs or steroid, drug addiction, history of cardiovascular, hepatic, respiratory, or renal diseases, presence of asthma or chronic pain syndrome

##### Age

From **35 years** old to **50 years** old

##### Gender

Female

## Phase

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

Kashan University of Medical Sciences

##### Street address

Vice-chancellor for Research, Ghotbe Ravandi Blvd, Kashan

##### City

Kashan

##### Postal code

#### Approval date

2010-03-16, 1388/12/25

#### Ethics committee reference number

7911

## Health conditions studied

### 1

#### Description of health condition studied

post abdominal hysterectomy pain

#### ICD-10 code

N99.9

#### ICD-10 code description

Postprocedural disorder of genitourinary system, unspecified

## Primary outcomes

### 1

#### Description

post-abdominal hysterectomy pain

#### Timepoint

2, 6, 12, 24 hours after operation

#### Method of measurement

Visual analog scale

## Secondary outcomes

### 1

#### Description

Morphine consumption

#### Timepoint

2, 6, 12, 24 hours after operation

#### Method of measurement

Number of morphine ampules consumption

## Intervention groups

### 1

#### Description

Gabapentine, 1200mg orally one hour before operation

#### Category

Treatment - Drugs

### 2

#### Description

Placebo, 1200mg orally similar to gabapentin exception for effective material one hour before operation

#### Category

Treatment - Drugs

## Recruitment centers

### 1

#### Recruitment center

##### Name of recruitment center

Shabih khani Maternity Hospital

##### Full name of responsible person

Dr. Fatemeh Foroozanfard

##### Street address

Shabih khani Maternity Hospital, Shahid Beheshti Street

##### City

Kashan

## Sponsors / Funding sources

### 1

#### Sponsor

##### Name of organization / entity

Vice-chancellor for Research, Kashan University of Medical Sciences

##### Full name of responsible person

Dr. Gholamali Hamidi

##### Street address

Vice-chancellor for Research, Ghotbe Ravandi

##### City

Kashan

#### 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, Kashan 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**

Kashan University of Medical Sciences

**Full name of responsible person**

Dr. Fatemeh Foroozanfard

**Position**

OB & GYN, IVF Fellowship, Assistant Professor

**Other areas of specialty/work****Street address**

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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*