

Clinical Trial Protocol

Iranian Registry of Clinical Trials

27 May 2026

Comparison of inguinal hernia recurrence rate in surgery by Liechtenstein method with bassini method with mesh embedded

Protocol summary

Summary

Aim of study: Comparison of inguinal hernia recurrence rate in surgery by Liechtenstein method with bassini method with mesh embedded. In a prospective clinical trial study, patients aged from 18 to 70 years who underwent unilateral inguinal hernia surgery at Velayat and Rajaei hospitals of Qazvin during years of 2015-2016, are entered to the study after obtaining informed consent. Exclusion criteria are as patients with history of hernia surgery, bilateral inguinal hernia, incarcerated hernia and femoral hernia. The patients are randomly (using closed envelopes and choose A or B cards) divided into two groups: the first group is underwent surgery by Liechtenstein technique with mesh; and the second group is repaired by basini technique. All patients refer for the first time for hernia repair surgery, and all of them are operated by a given surgical team. Postoperative pain intensity is evaluated based on Visual Analog Scale (VAS). Patients are followed for inguinal hernia recurrence immediately, 48 hours, 3 month, 6 month and one year after surgery by using phone call and questionnaire through clinic

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2017052834175N1**

Registration date: **2017-08-26, 1396/06/04**

Registration timing: **registered_while_recruiting**

Last update:

Update count: **0**

Registration date

2017-08-26, 1396/06/04

Registrant information

Name

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Name of organization / entity

Qazvin University of Medical Sciences

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Recruitment status

Recruitment complete

Funding source

Vice Chancellor for research, Qazvin University of Medical Sciences

Expected recruitment start date

2017-06-05, 1396/03/15

Expected recruitment end date

2017-09-21, 1396/06/30

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Comparison of inguinal hernia recurrence rate in surgery by Liechtenstein method with bassini method with mesh embedded

Public title

Comparison the recurrence of groin hernia surgery by two different surgical method.

Purpose

Prevention

Inclusion/Exclusion criteria

Inclusion criteria: unilateral inguinal hernia; candidate for surgery under general or spinal anesthesia. Exclusion criteria: patients younger than 18 and older than 70 years; patients with history of hernia surgery; bilateral

inguinal hernia; incarcerated hernia and femoral hernia.

Age

From **18 years** old to **70 years** old

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **80**

Randomization (investigator's opinion)

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

Ethical Comittee of Qazvin University of Medical Sciences

Street address

Qazvin, Shahid Bahonar Blvd

City

Qazvin

Postal code

Approval date

2017-04-08, 1396/01/19

Ethics committee reference number

IR.QUMS.REC.1396.59

Health conditions studied

1

Description of health condition studied

inguinal hernia, Surgical Wound Complications

ICD-10 code

K40.9

ICD-10 code description

Unilateral or unspecified inguinal hernia

2

Description of health condition studied

Complications of procedures

ICD-10 code

T81

ICD-10 code description

Complications of procedures, not elsewhere classified

Primary outcomes

1

Description

recurrence of inguinal hernia

Timepoint

48 hour, 3 month, 6 month and one year after surgery

Method of measurement

Examination

Secondary outcomes

1

Description

Surgical wound complications

Timepoint

48 hours, 3 month, 6 month and one year after surgery

Method of measurement

Examination

2

Description

Postoperative inguinal pain

Timepoint

immediately, 48 hours, 3 month, 6 month and one year after surgery

Method of measurement

Measurement of pain according to VAS scal

Intervention groups

1

Description

surgery by Liechtenstein technique

Category

Treatment - Surgery

2

Description

surgery by basini technique

Category

Treatment - Surgery

Recruitment centers

1

Recruitment center

Name of recruitment center

Velayat hospital

Full name of responsible person

Street address

City

Qazvin

2

Recruitment center

Name of recruitment center

Rajae hospital

Full name of responsible person

Street address

City

Qazvin

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Vice Chancellor for research, Qazvin University of
Medical Sciences

Full name of responsible person

Dr. Amir Peymani

Street address

Qazvin, Shahid Bahonar Blvd.

City

Qazvin

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, Qazvin 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

Qazvin University of Medical Sciences

Full name of responsible person

Dr. Nariman Mehrnia

Position

Assistant of Surgery

Other areas of specialty/work

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**Person responsible for scientific
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Contact

Name of organization / entity

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Associate professor of surgery

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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