

Clinical Trial Protocol

Iranian Registry of Clinical Trials

26 Jun 2026

Investigating the effect of magnetic therapy on the incidence of post-operative acute urinary retention in patients undergoing surgery in hospital: a randomized controlled trial

Protocol summary

Study aim

Determining the effect of magnetic therapy on the incidence of acute urinary retention after surgery in patients admitted to selected hospitals in Isfahan: a randomized controlled trial

Design

A total of 70 eligible patients are randomly divided into experimental and control groups as soon as they enter the recovery and will be followed up in the recovery and even in the surgical ward until the first urine is excreted after surgery. Will be affected by seven magnets. The total time the magnets are in place will be about 5.35 minutes. All stages of the intervention will be performed by the intervener who has been trained in acupuncture and magnetic therapy and will be coordinated and supervised by the treating physician. The length of time that the patient will be able to urinate after the installation of magnets or magnets is recorded in the experimental and control groups. Excreted urine volume will also be recorded.

Settings and conduct

This study is a single-blind clinical trial in the operating room recovery ward and in the men and women surgery ward of selected hospitals in Isfahan.

Participants/Inclusion and exclusion criteria

Patients at least 18 years old, without urinary catheterization, Class I and II according to the American Society of Anesthesiology (ASA) rankings, no drug addiction (according to the patient), no heart problems, elective and non-urological surgery

Intervention groups

In the experimental group, the magnets are glued at points P6, H7 on the right and left hands, and points SP6, ST36 only on the left foot, and a point on the sole of the left bladder / gastrointestinal foot. In the control group, they are placed at the same points of the facade without magnetic properties.

Main outcome variables

Elimination of urinary retention as a problem after surgery with the least cost and time and complications

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20210415050985N1**

Registration date: **2021-10-10, 1400/07/18**

Registration timing: **prospective**

Last update: **2021-10-10, 1400/07/18**

Update count: **0**

Registration date

2021-10-10, 1400/07/18

Registrant information

Name

Mohammad Sadegh Aboutalebi

Name of organization / entity

Country

Iran (Islamic Republic of)

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+98 31 3792 7574

Email address

aboutalebi@nm.mui.ac.ir

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2021-10-23, 1400/08/01

Expected recruitment end date

2022-02-20, 1400/12/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Investigating the effect of magnetic therapy on the incidence of post-operative acute urinary retention in patients undergoing surgery in hospital: a randomized controlled trial

Public title

The effect of magnetic therapy on the incidence of post-operative acute urinary retention in patients undergoing surgery in hospital

Purpose

Prevention

Inclusion/Exclusion criteria**Inclusion criteria:**

Patient informed consent Patients at least 18 years old Classes I and II are ranked according to the American Society of Anesthesiology (ASA) The duration of surgery was more than 2 hours Availability of places for putting magnets on the patient's body Elective surgeries Non-urological surgeries

Exclusion criteria:

Existence of urinary catheter Drug addiction Existence of heart problems History of previous urinary retention Existence of cardiovascular problems Existence of pacemaker Intervention to eliminate urinary retention and catheterization patient expressing a feeling of urination when entering to recovery

Age

From **18 years** old

Gender

Both

Phase

N/A

Groups that have been masked

- Participant

Sample size

Target sample size: **64**

Randomization (investigator's opinion)

Randomized

Randomization description

Sampling will be simple and accessible and according to the inclusion criteria. The allocation of samples in the control and test groups will be done randomly and individually using the website www.sealedenvelope.com in random mode. The samples will also be homogenized by minimization software in terms of age, sex and type of surgery and its duration.

Blinding (investigator's opinion)

Single blinded

Blinding description

The patient will not notice the magnet or non magnetic metal properties

Placebo

Not used

Assignment

Parallel

Other design features**Secondary Ids**

empty

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

Faculty of Nursing, Management and Rehabilitation - Isfahan University of Medical Sciences

Street address

Hezarjarib

City

Isfahan

Province

Isfahan

Postal code

8174993541

Approval date

2021-06-08, 1400/03/18

Ethics committee reference number

IR.MUI.NUREMA.REC.1400.047

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

urinary retention

ICD-10 code

R33

ICD-10 code description

Retention of urine

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

Elimination of urinary retention completely

Timepoint

During the study and the probable time of urination from the start of magnetite therapy

Method of measurement

obsevation

2**Description**

The degree of relaxation

Timepoint

Before placing the magnets and the time to remove them and one hour after removing them

Method of measurement

Orally asked based on visual numerical scale 0 to 10

3

Description

Nausea score

Timepoint

Before placing the magnets and the time to remove them and one hour after removing them

Method of measurement

Orally asked based on visual numerical scale 0 to 10

4

Description

Heart rate beat

Timepoint

Before placing the magnets and the time to remove them and one hour after removing them

Method of measurement

Digital monitor

5

Description

Average arterial blood pressure

Timepoint

Before placing the magnets and the time to remove them and one hour after removing them

Method of measurement

Barometer

Secondary outcomes

empty

Intervention groups

1

Description

Intervention group: Magnets are glued at points P6, H7 on the right and left hands, and points SP6, ST36 only on the left foot, and one point on the left foot of the bladder / gastrointestinal tract. The total time the magnets are in place will be about 35 minutes.

Category

Treatment - Other

2

Description

Control group: Non-magnetic irons are glued at points P6, H7 on the right and left hands, and points SP6, ST36 on the left foot only, and a point on the sole of the left foot corresponding to the bladder / gastrointestinal tract. The total time the magnets are in place will be about 5.35 minutes.

Category

Treatment - Other

Recruitment centers

1

Recruitment center**Name of recruitment center**

Amin hospital

Full name of responsible person

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Sponsors / Funding sources

1

Sponsor**Name of organization / entity**

Esfahan University of Medical Sciences

Full name of responsible person

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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

Esfahan 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

Name of organization / entity
Esfahan University of Medical Sciences
Full name of responsible person
Mohammad Sadegh Aboutalebi
Position
Consultant
Latest degree
Master
Other areas of specialty/work
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Person responsible for updating data

Contact

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Sharing plan

Deidentified Individual Participant Data Set (IPD)

Yes - There is a plan to make this available

Study Protocol

Yes - There is a plan to make this available

Statistical Analysis Plan

Yes - There is a plan to make this available

Informed Consent Form

Yes - There is a plan to make this available

Clinical Study Report

Yes - There is a plan to make this available

Analytic Code

Yes - There is a plan to make this available

Data Dictionary

Yes - There is a plan to make this available

Title and more details about the data/document

All data is potentially shareable after unidentified individuals

When the data will become available and for how long

2022

To whom data/document is available

It will be available only to researchers working in academic and scientific institutions

Under which criteria data/document could be used

no problem

From where data/document is obtainable

Mohammad Sadegh Aboutalebi

What processes are involved for a request to access data/document

As soon as possible

Comments