

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

09 Jun 2026

### Effect of vaginal misoprostol on cervical priming ic diagnostic dilatation and curettage

#### Protocol summary

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##### Summary

The purpose of the present study is to evaluate the effect of vaginal misoprostol on priming the cervix before dilatation in patients who are candidate for this procedure. Methods : A vandomized clinical trial was performed on 60 women who were candidated for dilation and curretage . In 30 patients ( case group), 200micro gram misoprostol ( one tablet ) was administered in posterior fornix of vagina 4 hours before operation , whereas in other 30 patients ( control group ), placebo was used. Then the two groups were compared according to their need to Hegar dilatator thinner than number 5 for dilatation of cervix and the duration of dilatation and curettage.

##### Recruitment status

**Recruitment complete**

##### Funding source

vice-chancellor for research , Qazvin university of medical sciences and heath services

##### Expected recruitment start date

2010-09-23, 1389/07/01

##### Expected recruitment end date

2020-01-21, 1398/11/01

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

#### General information

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT201110084868N3**

Registration date: **2011-12-20, 1390/09/29**

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

Last update:

Update count: **0**

##### Registration date

2011-12-20, 1390/09/29

##### Registrant information

###### Name

Khadijeh Elmizadeh

###### Name of organization / entity

Qazvin University of Medical Sciences

###### Country

Iran (Islamic Republic of)

###### Phone

+98 28 1333 6001

###### Email address

##### Scientific title

Effect of vaginal misoprostol on cervical priming ic diagnostic dilatation and curettage

##### Public title

use of vaginal misoprostol for cervical ripening

##### Purpose

Supportive

##### Inclusion/Exclusion criteria

Inclusion criteria :patints with abnormal uterine bleeding who are candidate for dilation and curretage exclusion criteria :systemic diseases ; cervical and vaginal infections; history of allergy to prostaglandins ; sever anemia or coagulopathy; anticoagulant therapy; history of cervical surgery;pregnancy; lactation; menopause; suspicious endocervical or exocervical erosions;

##### Age

From **27 years** old to **50 years** old

##### Gender

Female

##### Phase

2

## Groups that have been masked

No information

## Sample size

Target sample size: 60

## 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

Qazvin , university of medical sciences and health services

##### Street address

Qazvin

##### City

Qazvin

##### Postal code

#### Approval date

2010-09-22, 1389/06/31

#### Ethics committee reference number

28/20/4418

## Health conditions studied

### 1

#### Description of health condition studied

Diseases of the genitourinary system

#### ICD-10 code

N92

#### ICD-10 code description

Excessive, frequent and irregular menstruation

## Primary outcomes

### 1

#### Description

dilation of cervix as much as hegar dilator numbr five

#### Timepoint

four hour after using drug

#### Method of measurement

passage of hegar dilator numbr five

## Secondary outcomes

### 1

#### Description

Decrease in operation time

#### Timepoint

During operation

#### Method of measurement

Measurement with chronometer

## Intervention groups

### 1

#### Description

Intervention:30paitents misoprostol one tablet(200 micro gr)was inserted in the vagina 4 hours before surjery for one time.

#### Category

Treatment - Other

### 2

#### Description

Control :In 30paisents one placebo tablet was inserted in vagina 4hours befor surjery for one time.

#### Category

Placebo

## Recruitment centers

### 1

#### Recruitment center

##### Name of recruitment center

Kosar Haspital

##### Full name of responsible person

Dr Khadijeh Elmizadeh

##### Street address

, Taleghni stereet , Kosar Hospital, Qazvin City

##### City

Qazvin

## Sponsors / Funding sources

### 1

#### Sponsor

##### Name of organization / entity

vice- chancellor for research , Qazvin univercity of medical sciences and health services

##### Full name of responsible person

Dr Saeed Asefzadeh

##### Street address

Shahid Bahonar Blv , Qazvin City

##### 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 univercity of  
medical sciences and health services

**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 and health  
services

**Full name of responsible person**

Dr Khadijeh Elmizadeh

**Position**

Obstetrician and Ghyneecologist

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

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**Full name of responsible person**

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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*