

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

10 Jun 2026

### Comparison of vaginal misoprostol with Foley catheter for cervical ripening and labor induction

#### Protocol summary

##### Study aim

Comparison of mechanical and drug methods in preparing cervix

##### Design

In two parallel groups using Foley catheter and Prostaglandin

##### Settings and conduct

A randomized clinical trial is conducted on pregnant women referred to Besat Hospital in Sanandaj, who have an indication of the termination of pregnancy for any reason (due to delivery, post date pregnancy)

##### Participants/Inclusion and exclusion criteria

Women referring to the delivery block of Besat Hospital indicating the induction of labor. Inclusion criteria: Gestational age equal to or greater than 37 weeks; by ultrasonography of the first-trimester And last menstrual period, inappropriate cervix and Bishop score of 1 equal to or less than 4, single pregnancy, vortex display, intact chorionic membrane and satisfaction to participate in the research. Patients are also adjusting in gravid distribution. Patients with previous history of cesarean or previous surgery on the uterus, vaginal bleeding, or Placenta previa, or the possibility of early detachment of the placenta, regular uterine contractions and possible susceptibility to contraindications for use of prostaglandins, as well as Intrauterine growth restriction and severe preeclampsia, or problems in controlling fetal heart sounds, fetal distress, are excluded.

##### Intervention groups

Two groups which using prostaglandin and mechanical methods of Foley catheter

##### Main outcome variables

Tachysystole, umbilical cord prolapse, Meconium stain, Uterine atony, Neonate Apgar score, Bishop score, Labor speed, Induced safety, Induction failure

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20170326033142N2**  
Registration date: **2018-07-28, 1397/05/06**  
Registration timing: **retrospective**

Last update: **2018-07-28, 1397/05/06**

Update count: **0**

##### Registration date

2018-07-28, 1397/05/06

##### Registrant information

##### Name

sorayya rashid zadeh

##### Name of organization / entity

Kurdistan university of medical science

##### Country

Iran (Islamic Republic of)

##### Phone

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##### Email address

s.rashidzadeh@muk.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

UNIVERSITY OF MEDICAL SCIENCES KURDISTAN VICE  
CHANCELLOR IN RESEARCH AFFAIR

##### Expected recruitment start date

2016-09-22, 1395/07/01

##### Expected recruitment end date

2017-10-22, 1396/07/30

##### Actual recruitment start date

2016-09-22, 1395/07/01

##### Actual recruitment end date

2017-10-22, 1396/07/30

##### Trial completion date

empty

##### Scientific title

Comparison of vaginal misoprostol with Foley catheter for cervical ripening and labor induction

#### Public title

Traction effect with foley catheter on cervix ripening compared to vaginal misoprostol

#### Purpose

Treatment

#### Inclusion/Exclusion criteria

##### Inclusion criteria:

Women referring to the delivery block of Besat Hospital indicating the induction of labor. Gestational age equal to or greater than 37 weeks Satisfaction to participate in the research intact chorionic membrane Vortex presentation Single pregnancy Inappropriate cervix and Bishop score 1 equal to or less than 4

##### Exclusion criteria:

fetal distress Problems in controlling fetal heart sounds severe pre-eclampsia Intrauterine growth restriction Contraindications for the use of prostaglandins Regular uterine contractions Early placental detachment Placenta previa Vaginal bleeding Previous cesarean section or previous surgery on the uterus

#### Age

No age limit

#### Gender

Female

#### Phase

N/A

#### Groups that have been masked

No information

#### Sample size

Target sample size: **120**

Actual sample size reached: **120**

#### Randomization (investigator's opinion)

Randomized

#### Randomization description

Simple randomization

#### 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

KURDISTAN UNIVERSITY OF MEDICAL SCIENCES

###### Street address

Kurdistan University of Medical Sciences Vice Chancellor in Research Affair Pasdaran Ave.

###### City

Sanandaj

#### Province

Kurdistan

#### Postal code

6617713446

#### Approval date

2016-09-19, 1395/06/29

#### Ethics committee reference number

IR.MUK.REC.1396/105

### Health conditions studied

#### 1

##### Description of health condition studied

Cervical ripening

##### ICD-10 code

O80

##### ICD-10 code description

Encounter for full-term uncomplicated delivery

### Primary outcomes

#### 1

##### Description

Dilation

##### Timepoint

Every 1 hour

##### Method of measurement

Vaginal examination

#### 2

##### Description

Effacement

##### Timepoint

Every 1 hour

##### Method of measurement

Vaginal examination

#### 3

##### Description

Delivery progress time

##### Timepoint

Every 1 hour

##### Method of measurement

Vaginal examination

### Secondary outcomes

empty

### Intervention groups

#### 1

##### Description

In the first group (misoprostol group), misoprostol 25 µg (200µg Samisaz tablets divided by divaidier) is placed in the posterior fornix of the vagina and will be repeated

every 6 hours to a maximum of 3 doses (total 75 µg). If the patient does not spontaneously enter the phase of labor (regular uterine contractions accompanied by progressive changes in the cervix), after 12 hours of induced labor, oxytocin will be used to terminate the pregnancy.

**Category**

Treatment - Drugs

**2****Description**

In the second group (Foley catheter group), Foley catheter No. 16 in sterile conditions (washing the vagina with Betadine), through the cervix into the uterus and then the balloon is filled with 30 ml of distilled water to be placed behind the inner hole To put The other end of the catheter, to provide traction with normal saline 500ml bag and hang beside the bed. If after 6 hours, the patient does not spontaneously enter the phase of labor, induction of labor with oxytocin begins.

**Category**

Treatment - Other

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

Be'sat Hospital, Sanandaj

**Full name of responsible person**

Dr. Sorayya Rashid Zadeh

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Be'sat Educational Hospita, Keshavarz St. Vakil 4way, sanandaj, Kurdistan, Iran

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

Sanandaj University of Medical Sciences

**Full name of responsible person**

Dr.Nasrin Soufi Zadeh

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**Grant name**

پایان نامه دستیار تخصصی جراح زنان و زایمان

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

Yes

**Title of funding source**

Sanandaj 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**

Sanandaj University of Medical Sciences

**Full name of responsible person**

Dr. Sorayya Rashid Zadeh

**Position**

Obstetrics and Gynecology Resident

**Latest degree**

Medical doctor

**Other areas of specialty/work**

Gynecology and Obstetrics

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**Web page address****Person responsible for scientific inquiries****Contact****Name of organization / entity**

Sanandaj University of Medical Sciences  
**Full name of responsible person**  
Dr. Shoaleh Shahgheybi  
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Specialist  
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## Person responsible for updating data

### Contact

**Name of organization / entity**  
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**Full name of responsible person**  
Behzad Khalafi  
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Research Associate, Researcher Soldier, General  
Physician of Kurdistan University of Medical Scien  
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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

Not applicable

### 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

Not applicable

### Data Dictionary

Not applicable

### Title and more details about the data/document

All data is from the proposal, raw data and project reports

### When the data will become available and for how long

Data is available from the legal system six months after publication for two years

### To whom data/document is available

All persons will be able to access the Kurdistan University of Medical Sciences's request to the Kurdistan University of Medical Sciences.

### Under which criteria data/document could be used

For legal issues and the need to use data in future studies

### From where data/document is obtainable

Sorayya Rashidzadeh

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

After submitting a request to the Kurdistan Research and Technology Dept. of Science and Technology, a call for submission is given.

### Comments