

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

19 Jun 2026

### A comparative study on the prophylactic effect of Dexmedetomidine and Paracetamol on intraoperative hemodynamic control and postoperative pain in patients undergoing laparoscopic cholecystectomy

#### Protocol summary

##### Study aim

Comparison of prophylactic effect of Dexmedetomidine and Paracetamol on hemodynamic control during surgery and post operative pain in patients with laparoscopic cholecystectomy

##### Design

Two arm parallel group randomized trial

##### Settings and conduct

132 patients are divided into two groups of 66 people. Patients admitted to the Amirmomenin Hospital of Arak are under study. In each group, hemodynamic therapy is measured before surgery, 30, 60, and 90 minutes postoperatively. Postoperative pain is measured by Visual Analogue Scale in recovery, 4, 12, and 24 hours postoperatively.

##### Participants/Inclusion and exclusion criteria

Inclusion criteria: 1: Patients from 18 to 70 years old 2: Both genders 3: Patients who are indicated for urgent or elective Laparoscopic Cholecystectomy 4: Acquiring the informed consent of participating in a clinical study  
Exclusion criteria: 1: Conversion from Laparoscopic to open surgery 2: Partial Cholecystectomy 3: Gangrenous Gall Bladder 4: Abscess 5: Trauma to major Bile ducts

##### Intervention groups

The Dexmedetomidine group, in which patients receive Dexmedetomidine before and after the onset of action. The Paracetamol group in which patients receive Paracetamol before and 6 hours to 24 hours.

##### Main outcome variables

Operative hemodynamics in which the MAP and heart rate are measured. Postoperative pain

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20180115038381N1**

Registration date: **2019-02-18, 1397/11/29**

Registration timing: **retrospective**

Last update: **2019-02-18, 1397/11/29**

Update count: **0**

##### Registration date

2019-02-18, 1397/11/29

##### Registrant information

###### Name

Taha Hojjati

###### Name of organization / entity

###### Country

Iran (Islamic Republic of)

###### Phone

+98 86 3223 1102

###### Email address

taha.hojjati@arakmu.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2017-05-05, 1396/02/15

##### Expected recruitment end date

2018-02-20, 1396/12/01

##### Actual recruitment start date

2017-04-04, 1396/01/15

##### Actual recruitment end date

2017-11-22, 1396/09/01

##### Trial completion date

2017-12-22, 1396/10/01

##### Scientific title

A comparative study on the prophylactic effect of Dexmedetomidine and Paracetamol on intraoperative hemodynamic control and postoperative pain in patients

undergoing laparoscopic cholecystectomy

## Public title

A comparative study on the prophylactic effect of Dexmedetomidine and Paracetamol on intraoperative hemodynamic control and postoperative pain in patients undergoing laparoscopic cholecystectomy

## Purpose

Treatment

## Inclusion/Exclusion criteria

### Inclusion criteria:

Patients who are indicated for emergency or elective Laparoscopic Cholecystectomy. Obtaining informed consent Both genders Patients between 18 to 70 years old

### Exclusion criteria:

Conversion of Laparoscopic surgery to open surgery  
Partial Cholecystectomy Gangrenous Gall bladder  
Abscess Major trauma to Bile ducts

## Age

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

## Gender

Both

## Phase

3

## Groups that have been masked

*No information*

## Sample size

Target sample size: **132**

Actual sample size reached: **132**

## Randomization (investigator's opinion)

Randomized

## Randomization description

Blocked Randomization Individual random unit  
Randomization tool under web ([www.randomizer.org](http://www.randomizer.org))  
Randomization sequence was made under web ([www.randomizer.org](http://www.randomizer.org)) Hiding: Sealed envelope

## 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

Ethics committee of Arak University of Medical Sciences

##### Street address

Deputy Director of Research and Technology,  
University of Medical Sciences campus site, Basij  
Square, Arak

##### City

Arak

## Province

Markazi

## Postal code

3819693345

## Approval date

2017-05-01, 1396/02/11

## Ethics committee reference number

IR.ARAKMU.REC.1396.1

## Health conditions studied

### 1

#### Description of health condition studied

Postoperative pain after Laparoscopic Cholecystectomy

#### ICD-10 code

#### ICD-10 code description

### 2

#### Description of health condition studied

Peri-operative Hemodynamic

#### ICD-10 code

#### ICD-10 code description

## Primary outcomes

### 1

#### Description

Mean Arterial Blood Pressure

#### Timepoint

Immediately after Anesthesia induction, 30 minutes after surgery, 1 hour after surgery, 90 minutes after surgery

#### Method of measurement

Blood pressure gauge

### 2

#### Description

Pulse Rate

#### Timepoint

Immediately after Anesthesia induction, 30 minutes after surgery, 1 hour after surgery, 90 minutes after surgery

#### Method of measurement

Measured by the questioner

### 3

#### Description

Postoperative Pain

#### Timepoint

In Recovery Unit, 4 hour after surgery, 12 hour after surgery, 24 hour after surgery

#### Method of measurement

With Visual Analogue Scale

## Secondary outcomes

empty

## Intervention groups

### 1

#### Description

The first intervention group included 66 patients undergoing laparoscopic cholecystectomy and receiving doxedemotomidine. Patients in this group receive 1 µg / Kg stat and then 0.5 µg / Kg per hour. In this study, the American Percedex product, each vial containing 200 µg doxedetomidine, is used.

#### Category

Treatment - Drugs

### 2

#### Description

Second intervention group: includes 66 patients undergoing laparoscopic cholecystectomy and receiving paracetamol. These patients first receive 1 g of intravenous paracetamol before surgery and receive 500 mg paracetamol every 6 hours after surgery.

#### Category

Treatment - Drugs

## Recruitment centers

### 1

#### Recruitment center

##### Name of recruitment center

Amiralmomenin Hospital

##### Full name of responsible person

Taha Hojjati Ashrafi

##### Street address

Basig square

##### City

Arak

##### Province

Markazi

##### Postal code

3848176941

##### Phone

+98 86 3417 3601

##### Fax

+98 86 3417 3619

##### Email

it-amiralmomenin@arakmu.ac.ir

## Sponsors / Funding sources

### 1

#### Sponsor

##### Name of organization / entity

Arak University of Medical Sciences

##### Full name of responsible person

Dr. Mohammad Arjomandzadegan

##### Street address

Basij Square,

##### City

Arak

#### Province

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#### Postal code

3848176341

#### Phone

+98 86 3417 3645

#### Email

mmatinam81@yahoo.com

#### Grant name

#### Grant code / Reference number

#### Is the source of funding the same sponsor organization/entity?

Yes

#### Title of funding source

Arak 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

Arak University of Medical Sciences

##### Full name of responsible person

Taha Hojjati Ashrafi

##### Position

Surgery Resident

##### Latest degree

Medical doctor

##### Other areas of specialty/work

General Surgery

##### Street address

4, Zeytoon Appartment, Behdari Street

##### City

Arak

##### Province

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3819693357

##### Phone

+98 86 3313 1639

##### Email

taha.hojjati@arakmu.ac.ir

## Person responsible for scientific inquiries

#### Contact

##### Name of organization / entity

Arak University of Medical Sciences

##### Full name of responsible person

Taha Hojjati Ashrafi

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Surgery Resident  
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## Person responsible for updating data

**Contact**  
**Name of organization / entity**  
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**Full name of responsible person**  
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Surgery Resident  
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## Sharing plan

**Deidentified Individual Participant Data Set (IPD)**  
Undecided - It is not yet known if there will be a plan to make this available  
**Study Protocol**  
Undecided - It is not yet known if there will be a plan to make this available  
**Statistical Analysis Plan**  
Undecided - It is not yet known if there will be a plan to make this available  
**Informed Consent Form**  
Undecided - It is not yet known if there will be a plan to make this available  
**Clinical Study Report**  
Undecided - It is not yet known if there will be a plan to make this available  
**Analytic Code**  
Undecided - It is not yet known if there will be a plan to make this available  
**Data Dictionary**  
Undecided - It is not yet known if there will be a plan to make this available