

Clinical Trial Protocol

Iranian Registry of Clinical Trials

20 Jul 2026

Comparison of analgesic effect of remifentanyl with and without dexamethasone in labor

Protocol summary

Study aim

Comparison of analgesic effect of remifentanyl with and without dexamethasone in labor

Design

In this interventional study, 100 pregnant women who are candidates for natural childbirth will be selected and divided into two groups using block randomization method (n=50 in each group). Proper counseling will be done and a written informed consent will be obtained before starting the treatment regimen.

Settings and conduct

Upon starting the active phase of labor intervention group will receive intravenous Remifentanyl 0.05 µg/kg/min (Tofigh Daru Co.) and 8 mg intravenous Dexamethasone (once, 2 cc, Irandaroo Co.) injection and control group receive intravenous Remifentanyl 0.05 µg/kg/min (Tofigh Daru Co.) and placebo (distilled water, 2 cc, Irandaroo Co., once) injection.

Participants/Inclusion and exclusion criteria

Inclusion criteria: Over 37 weeks pregnancy, first or second pregnancy, singleton. Exclusion criteria: Malpresentation of fetus, macrosomia.

Intervention groups

Intravenous Remifentanyl 0/05 µg/kg/min (Tofigh Daru Co.) and 8 mg intravenous Dexamethasone (once, 2 cc, Irandaroo Co.) injection. Control group: Intravenous Remifentanyl 0/05 µg/kg/min (Tofigh Daru Co.) and placebo (distilled water, 2 cc, Irandaroo Co., once) injection.

Main outcome variables

pain

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20150808023559N16**

Registration date: **2018-03-10, 1396/12/19**

Registration timing: **prospective**

Last update: **2018-03-10, 1396/12/19**

Update count: **0**

Registration date

2018-03-10, 1396/12/19

Registrant information

Name

Somaieh Matin

Name of organization / entity

Ardabil University of Medicine Sciences

Country

Iran (Islamic Republic of)

Phone

+98 45 3373 3011

Email address

s.matin@arums.ac.ir

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2018-04-19, 1397/01/30

Expected recruitment end date

2018-06-20, 1397/03/30

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Comparison of analgesic effect of remifentanyl with and without dexamethasone in labor

Public title

Labor pain relief with remifentanyl

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria:
Over 37 weeks pregnancy First or second pregnancy
Singleton

Exclusion criteria:
Malpresentation of fetus Macrosomia

Age
No age limit

Gender
Female

Phase
2

Groups that have been masked

- Participant
- Care provider

Sample size
Target sample size: **100**

Randomization (investigator's opinion)
Randomized

Randomization description
Patients are assigned randomly by blocking in one of the two specified groups and after selecting the envelope for each patient, before starting the treatment regimen envelope will be opened and based on the protocol, one of two methods will be selected.

Blinding (investigator's opinion)
Double blinded

Blinding description
Participants and clinical caregivers are not aware of the type of medication

Placebo
Used

Assignment
Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee
Ethics committee of Ardabil University of Medical Sciences

Street address
Ardabil University of Medicine Sciences, Daneshgah street, Ardabil

City
Ardabil

Province
Ardabil

Postal code
5615783134

Approval date
2017-12-31, 1396/10/10

Ethics committee reference number
IR.ARUMS.REC.1396.189

Health conditions studied

1

Description of health condition studied
Labor pain

ICD-10 code
O80, O84

ICD-10 code description
Delivery

Primary outcomes

1

Description
pain

Timepoint
Before analgesic administrations and two times (30 and 60 min) after intervention

Method of measurement
Visual analog scale

Secondary outcomes

1

Description
Apgar

Timepoint
1 and 5 minutes after birth

Method of measurement
Apgar test

Intervention groups

1

Description
Intervention group: Intravenous Remifentanil 0/05 µg/kg/min (Tofigh Daru Co.) and 8 mg intravenous Dexamethasone (once, 2 cc, Irandaroo Co.) injection

Category
Treatment - Drugs

2

Description
Control group: Intravenous Remifentanil 0/05 µg/kg/min (Tofigh Daru Co.) and placebo (distilled water, 2 cc, Irandaroo Co., once) injection

Category
Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Alavi Hospital

Full name of responsible person

Somaieh Sagafi rad

Street address

Alavi Hospital, Moadi street, Ardabil

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5615783134

Phone

+98 45 4233 8888

Email

drssrad62@gmail.com

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Ardabil University of Medical Sciences

Full name of responsible person

Masoud Entezari

Street address

Ardabil University of Medical Science, Daneshgah street, Ardabil, Iran

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m.entezari@arums.ac.ir

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

Yes

Title of funding source

Ardabil 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

Ardabil University of Medical Sciences

Full name of responsible person

Somaieh Sagafi rad

Position

Gynecology Asisstant

Latest degree

Medical doctor

Other areas of specialty/work

Gynecology and Obstetrics

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Person responsible for scientific inquiries

Contact

Name of organization / entity

Ardabil University of Medical Sciences

Full name of responsible person

Noshin Mobaraki

Position

Associate professor

Latest degree

Specialist

Other areas of specialty/work

Gynecology and Obstetrics

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Person responsible for updating data

Contact

Name of organization / entity

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Full name of responsible person

Somaieh Sagafi rad

Position

Gynecology Asisstant

Latest degree

Medical doctor

Other areas of specialty/work

Gynecology and Obstetrics

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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