Effect of combined Paracetamol(Apotel) and Dexamethasone versus Paracetamol(Apotel) on post operative nausea vomiting after cesarean section

Protocol summary

Study aim
Effect of combined Paracetamol(Apotel) and Dexamethasone versus Paracetamol(Apotel) on post operative nausea vomiting after cesarean section

Design
Clinical trial with 2 parallel groups, double blind, randomised, phase 3 on 100 patients, in block randomization method with the size of 4 and 6. Random sequence will be generated by running an online program in sealed envelope website (https://www.sealedenvelope.com/).

Settings and conduct
This is a double blind clinical trial study with 100 patients referred to Taleghani hospital.They will be devided into 2 groups (Apotel and combined Apotel-Dexamethasone)randomly.

Participants/Inclusion and exclusion criteria
Inclusion criteria: Age between 18-45 years Having informed consent Having ASA I and II (American Society of Anesthesiologists) Nulliparous Singleton pregnancy Patients with elective cesarean Patients with spinal anesthesia Patients with Pfannenstiel incision in surgery. Non-inclusion criteria: Underlying disease (i.e renal, liver, cardiac and pulmonary disease) History of allergy to the drugs of study Addiction or long term using of opiate Coagulopathy or use of anticoagulant drugs Disability for pain scoring Suffering from preeclampsia or HELLP (hemolysis,elevated liver enzymes,low platelet count) or eclampsia or hepatitis Urgency cesarean section

Intervention groups
In first group 2 cc of distilled water and in second group 8 mg of Dexamethasone 30 minutes before spinal anesthesia and in both groups 1 gr of Apotel immediately after spinal anesthesia will be intravenously injected to the patients.

Main outcome variables
Vomiting score and happening of nauseae after cesarean section till 24 hours

General information

Reason for update
Acronym

IRCT registration information
IRCT registration number: IRCT20201028049175N6
Registration date: 2021-03-02, 1399/12/12
Registration timing: registered_while_recruiting

Last update: 2021-03-02, 1399/12/12
Update count: 0

Registration date
2021-03-02, 1399/12/12

Registrant information
Name
Shamim Valibak
Name of organization / entity
Country
Iran (Islamic Republic of)
Phone
+98 86 3417 3502
Email address
sh.valibak@yahoo.com

Recruitment status
Recruitment complete
Funding source

Expected recruitment start date
2021-02-19, 1399/12/01
Expected recruitment end date
2021-08-23, 1400/06/01
Actual recruitment start date
empty
Actual recruitment end date
empty
Trial completion date
empty

Scientific title
Effect of combined Paracetamol(Apotel) and Dexamethasone versus Paracetamol(Apotel) on post operative nausea vomiting after cesarean section

Public title
Evaluation of anti analgesic and anti inflammatory drugs effect on decreasing of post operative nausea vomiting after cesarean section

Purpose
Prevention

Inclusion/Exclusion criteria
Inclusion criteria:
Age between 18-45 years Having informed consent Having ASA I and II Nulliparous Singleton pregnancy Patients with elective cesarean Patients with spinal anesthesia Patients with Pfannenstiel incision in surgery.

Exclusion criteria:
Underlying disease (i.e renal, liver, cardiac and pulmonary disease) History of allergy to the drugs of study Addiction or long term using of opiate Coagulopathy or use of anticoagulant drugs Disability for pain scoring Suffering from preeclampsia or HELLP or eclampsia or hepatitis Urgency cesarean section

Age
From 18 years old to 45 years old

Gender
Female

Phase
3

Groups that have been masked
• Participant
• Investigator

Sample size
Target sample size: 100

Randomization (investigator's opinion)
Randomized

Randomization description
To create simple random sequences web randomization (com.graphpad.www) dual group and for concealment, closed envelope method with random sequence (SNOSE) will be used.Random sequences will be recorded on the cards and then sequentially puted in the envelopes.In order to preserve random sequences the outer surface of the envelopes are numbered, respectively. Finally, the lid of the letter envelopes was pasted and it is placed inside the boxes, respectively.At the time of participants registration, one of the envelopes will be opened and the allocation group, will be revealed.

Blinding (investigator's opinion)
Double blinded

Blinding description
Patients blindness: According to injecting of Dexamethasone in one group of study 30 minutes before spinal anesthesia, in another group distilled water will be injected to the patients in same time and measure. Researcher blindness: Drugs will be ordered by specialist, so the researcher who filled the check lists is blind.

Placebo
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
Assistance of research and technology, Payambare azam institution, Arak University of Medical Sciences, Basij square

City
Arak

Province
Markazi

Postal code
3848176341

Approval date
2021-01-03, 1399/10/14

Ethics committee reference number
IR.ARAKMU.REC.1399.290

Health conditions studied

1

Description of health condition studied
Nausea and vomiting after cesarean

ICD-10 code
R11

ICD-10 code description
Nausea and vomiting

Primary outcomes

1

Description
Nausea happening after cesarean section until 24 hours after surgery

Timepoint
In recovery room and 2,4,6,12,24 hours after surgery

Method of measurement
Asking of patient

2

Description
Vomiting score after cesarean section until 24 hours after surgery
**Timepoint**  
In recovery room and 2,4,6,12,24 hours after surgery

**Method of measurement**  
VAS (Visual Analog Scale) score

**Secondary outcomes**  
empty

**Intervention groups**

1

**Description**  
Intervention group: 8 mg of Dexamethasone (Daroo pakhsh company) intravenously will be injected to the patients, 30 minutes before spinal anesthesia and 1 gr of Apotel (Exir company) intravenously will be injected immediately after spinal anesthesia.

**Category**  
Treatment - Drugs

2

**Description**  
Control group: 2 cc of distilled water intravenously will be injected to the patients, 30 minutes before spinal anesthesia and 1 gr of Apotel (Exir company) intravenously will be injected immediately after spinal anesthesia.

**Category**  
Treatment - Drugs

**Recruitment centers**

1

**Recruitment center**  
Name of recruitment center  
Taleghani hospital  
Full name of responsible person  
Dr Maryam Maktabi  
Street address  
West side of Emam Khomeini street, beside of gas company  
City  
Arak  
Province  
Markazi  
Postal code  
3816149369  
Phone  
+98 86 3277 6063  
Email  
it-taleghani@arakmu.ac.ir

**Sponsors / Funding sources**

1

**Sponsor**  
Name of organization / entity  
Arak University of Medical Sciences  
Full name of responsible person  
Alireza Kamali  
Street address  
Assistance of research and technology, Payambarazam institute, Arak University of Medical Sciences, Basij square  
City  
Arak  
Province  
Markazi  
Postal code  
3848176341  
Phone  
+98 86 3417 3639  
Email  
research@arakmu.ac.ir  
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  
Maryam Maktabi  
Position  
Assistant professor  
Latest degree  
Subspecialist  
Other areas of specialty/work  
Gynecology and Obstetrics  
Street address  
West side of Emam Khomeini street, beside of gas company  
City  
Arak  
Province  
Markazi  
Postal code  
3816149369  
Phone  
+98 86 3277 6063  
Email  
Dr.maryam.maktabi@gmail.com
Person responsible for scientific inquiries

Contact
Name of organization / entity
Arak University of Medical Sciences
Full name of responsible person
Alireza Kamali
Position
Associate Professor
Latest degree
Specialist
Other areas of specialty/work
Anesthesiology
Street address
Valiasr hospital, Valiasr square
City
Arak
Province
Markazi
Postal code
3814957558
Phone
086-32222003-8
Email
alikamaliir@yahoo.com

Person responsible for updating data

Contact
Name of organization / entity
Arak University of Medical Sciences
Full name of responsible person
Shamim Valibak
Position
General physician non-faculty
Latest degree
Medical doctor

Other areas of specialty/work
General Practitioner
Street address
Valiasr hospital, Valiasr square
City
Arak
Province
Markazi
Postal code
3814957558
Phone
086-32222003-8
Email
sh.valibak@yahoo.com

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