

Clinical Trial Protocol

Iranian Registry of Clinical Trials

04 Jul 2026

The effect of preoperative dextrose infusion on postoperative nausea and vomiting

Protocol summary

Summary

Postoperative nausea and vomiting (PONV) remains one of the most common postoperative complications. The aim of the present study is to determine the Efficacy of preoperative dextrose infusion on postoperative nausea and vomiting. In this clinical trial 300 ASA physical status I non-diabetic patients and aged 18-40 years, scheduled for open surgeries are randomly assigned to infusion of 10 ml/kg/h Ringer's solution with 100 mL/h dextrose 10% in study group (n = 150) or Ringer's solution alone in placebo group (n = 150) during surgery. After establishment of monitoring of the operating room, all patients will be placed under general anesthesia with the same anesthetic technique, including midazolam 0.01 mg / kg, fentanyl 2µg / kg for Pre-medication, Propofol 2-2.5 mg / kg, cisatracurium 0.15 mg / kg for induction of anesthesia and isoflurane 1-1.5 MAC, morphine 0.1 mg / kg and cisatracurium during maintenance of anesthesia. PONV scores using by Visual Analogue Scale (VAS) and (Nausea and vomiting in slight level is zero to 3, in average level is 3 to 7 and in severe level is more than 7) scoring in recovery room and at 0, 30, 60, 120 minute and 4 hours postoperative are recorded.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2015032721544N1**
Registration date: **2015-06-21, 1394/03/31**
Registration timing: **registered_while_recruiting**

Last update:

Update count: **0**

Registration date

2015-06-21, 1394/03/31

Registrant information

Name

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Name of organization / entity

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Recruitment status

Recruitment complete

Funding source

Vice chancellor for research, Shahroud University of Medical Sciences

Expected recruitment start date

2015-06-21, 1394/03/31

Expected recruitment end date

2015-11-21, 1394/08/30

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The effect of preoperative dextrose infusion on postoperative nausea and vomiting

Public title

The effect of dextrose infusion on postoperative nausea and vomiting

Purpose

Prevention

Inclusion/Exclusion criteria

Inclusion criteria: ASA class 1; both sexes; aged 18 to 40 years; Open surgeries that are identical in terms of time and type of anesthesia. Exclusion criteria: diabetics; history of systemic disease (cardiovascular, respiratory,

renal ,...); history of nausea and vomiting.

Age

From **18 years** old to **40 years** old

Gender

Both

Phase

3

Groups that have been masked

No information

Sample size

Target sample size: **298**

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

Ethics committee, Vice chancellor for research,
Shahroud University of Medical Sciences

Street address

Ethics committee, Vice chancellor for research,
Shahroud University of Medical Sciences, 7th of Tir
square, Shahroud, Iran. Postal code: 3614773955

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Approval date

2015-05-13, 1394/02/23

Ethics committee reference number

IR.SHMU.REC.1394.17

Health conditions studied

1

Description of health condition studied

Postoperative Nausea and Vomiting

ICD-10 code

Y84

ICD-10 code description

Other medical procedures as the cause of abnormal
reaction of the patient, or of later complication, without
mention of misadventure at the time of the procedure

Primary outcomes

1

Description

Intensity of Postoperative Nausea and Vomiting

Timepoint

0 (while the patient is awake), 30, 60, 120 minute and 4
hours after the surgery.

Method of measurement

Questionnaire- Visual Analog Scale

Secondary outcomes

1

Description

Total dose antiemetic drug

Timepoint

Recovery within 24 hours after surgery

Method of measurement

Questionnaire

Intervention groups

1

Description

In the study group (n=150) is infused 10 ml/kg/h Ringer's
solution with 100 mL/h dextrose 10% during surgery.

Category

Prevention

2

Description

In the placebo group (n=150) is infused of 10 ml/kg/h
Ringer's solution alone in placebo during surgery.

Category

Placebo

Recruitment centers

1

Recruitment center

Name of recruitment center

Operation room, Imam Hossein hospital

Full name of responsible person

Javad Nourian MD

Street address

Imam Hossein Hospital, operation room, End of
Towhidi avenue

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Vice chancellor for research, Shahroud University of Medical Sciences

Full name of responsible person

Mohammad Hassan Emamian MD

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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, Shahroud University of Medical Sciences

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**

Imam Hossein Hospital, Shahroud University of Medical Sciences, Shahroud, Iran

Full name of responsible person

Javad Nourian MD

Position

Assistant professor of anesthesiology, department of anesthesia and intensive care

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Operating Room student

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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