

Clinical Trial Protocol

Iranian Registry of Clinical Trials

10 Jun 2026

The Effect Comparison of synchronized intermittent mandatory ventilation and airway pressure release ventilation in oxygation and hemodynamic parameters in patients with head trauma.

Protocol summary

Summary

Objective: To determine the effect of APRV on oxygation and hemodynamic of patients with traumatic brain injuries. This study is a clinical trial with a crossover design and subjects will be allocated into two groups according to the table of random numbers. Study population: patients with traumatic brain injury. Inclusion criteria: age between 15 and 60 years; patients with traumatic brain injury who are intubated and are under mechanical ventilation with SIMV mode; stable hemodynamic (MAP > 70 mm Hg); Glasgow Coma Scale <9. Exclusion criteria: unstable hemodynamic (MAP <70), need to CPR. Sample size: 36 patients in each group. Intervention of this research is a ventilation mode called Airway Pressure Release Ventilation that will be applied using ventilator and tracheal tube for half an hour. This mode is an inverse ratio mode in which patient airway pressure is fluctuated between high pressure and low pressure. In this intervention, the duration of high pressure is 4.5 sec and duration of low pressure is 0.5 sec. The high-level pressure will be determined based on the patient's required tidal volume. All patients will receive synchronized intermittent mandatory ventilation (SIMV). Then the ventilation mode in patients in group I will be switched to APRV while in group II to SIMV and will be continue for another two hours. Afterwards, group I will receive SIMV and group II will receive APRV for two hours. In all three stages the values for ABG and vital signs (pao₂, paco₂, spo₂, MAP, heart rate, blood pressure) will be recorded.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2016080729234N1**
Registration date: **2017-06-06, 1396/03/16**

Registration timing: **retrospective**

Last update:

Update count: **0**

Registration date

2017-06-06, 1396/03/16

Registrant information

Name

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Name of organization / entity

Sabzevar University of Medical Sciences

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Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Vice chancellor for research, Sabzevar University of Medical Sciences .

Expected recruitment start date

2016-08-22, 1395/06/01

Expected recruitment end date

2016-12-20, 1395/09/30

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The Effect Comparison of synchronized intermittent mandatory ventilation and airway pressure release

ventilation in oxynation and hemodynamic parameters in patients with head trauma.

Public title

Effect of mechanical ventilation on oxynation and hemodynamic in head trauma.

Purpose

Supportive

Inclusion/Exclusion criteria

Inclusion criteria: age between 15 and 60 years; patients suffering from traumatic brain injuries who are intubated and are under mechanical ventilation with SIMV mode; stable hemodynamic (MAP more than 70 mm Hg for at least 30 minutes before the start of the study); Glasgow Coma Scale <9 ; no history of chronic obstructive pulmonary disease (COPD); Exclusion criteria: unstable hemodynamic (MAP <70); need to CPR.

Age

From **15 years** old to **60 years** old

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **72**

Randomization (investigator's opinion)

Randomized

Randomization description**Blinding (investigator's opinion)**

Not blinded

Blinding description**Placebo**

Not used

Assignment

Other

Other design features**Secondary Ids**

empty

Ethics committees**1****Ethics committee****Name of ethics committee**

Ethics committee of Sabzevar University of Medical Sciences.

Street address

Central beuilding of Sabzevar University of Medical Sciences, Assadabadi street, Sabzevar.

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Sabzevar

Postal code

9613873136

Approval date

2016-12-01, 1395/09/11

Ethics committee reference number

IR.MEDSAB.REC.1395.103

Health conditions studied**1****Description of health condition studied**

Head trauma

ICD-10 code

s06.2, s06

ICD-10 code description

Other intracranial injuries, Intracranial injury with prolonged coma, Traumatic subarachnoid haemorrhage, Traumatic subdural haemorrhage, Diffuse brain injury, Focal brain injury, Epidural haemorrhage,

Primary outcomes**1****Description**

Pressure of arterial oxygen

Timepoint

Before intervention, half an hour and 2 hours after intervention.

Method of measurement

Atrial blood gas (ABG).

2**Description**

Pressure of arterial CO2

Timepoint

Before intervention, half an hour and 2 hours after intervention.

Method of measurement

Atrial blood gas (ABG)

3**Description**

Aterial oxygen saturation

Timepoint

Before intervention, half an hour and 2 hours after intervention.

Method of measurement

Monitore

Secondary outcomes**1****Description**

Mean artery pressure

Timepoint

Before intervention, half an hour and 2 hours after intervention.

Method of measurement

Sphygmomanometer

2**Description**

Heart rate

Timepoint

Before intervention, half an hour and 2 hours after intervention.

Method of measurement

Number in minute

3**Description**

Systolic pressure

Timepoint

Before intervention, half an hour and 2 hours after intervention.

Method of measurement

Sphygmomanometer

Intervention groups**1****Description**

Intervention of this research is a ventilation mode is called Airway Pressure Release Ventilation that will be applied through ventilator and tracheal tube for half an hour. This mode is an inverse ratio mode in which patient airway pressure is fluctuated between high pressure and low pressure. In this intervention, the duration of high pressure is 4.5 sec and duration of low pressure is 0.5 sec. The high-level pressure will be determined based on the patient's required tidal volume. The patients in group 1 will receive SIMV for two hours then ABG will be taken half an hour and two hours after changes. Then ventilator mode will be switched to APRV for two hours and ABG will be taken half an hour and two hours later. In all stages the values for ABG and vital signs will be recorded.

Category

Other

2**Description**

The patients in group 2 will receive APRV for two hours and after that ABG will be taken half an hour and two hours after changes. Then ventilator mode will be switched to SIMV for two hours and ABG will be taken half an hour and two hours later. In all stages the values for ABG and vital signs will be recorded.

Category

Other

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

Mashhad Taleghani Hospita

Full name of responsible person

Dr Hassan Sadeghi

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

Vice Chancellor for research of Sabzevar University of Medical Sciences

Full name of responsible person

Akrami Rahim

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City

Sabzevar

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 of Sabzevar 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**

Sabzevar University of Medical Sciences

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Position

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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