

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

26 Jun 2026

### **Determining and comparing the effect of Adaptive Support Ventilation and Synchronized Intermittent Mandatory Ventilation on respiratory support outcomes after Coronary Artery Bypass Graft surgery, Isfahan University of Medical sciences, Chamran hospital, 2014**

#### **Protocol summary**

##### **Summary**

Aim: Comparing the effect(s) of using adaptive support ventilation and synchronized intermittent mandatory ventilation on respiratory support outcomes (ventilation duration, length of hospital stay, unsuccessful weaning cases, and haemodynamic parameters of mean atrial blood pressure (MAP) and SPO<sub>2</sub> ) after coronary artery bypass graft surgery. Design: Randomized, one period, single blinded, two groups Population: Patients after coronary artery bypass graft surgery which were transferred to cardiac surgery ICU Main inclusion criteria: The range of age was between 25 and 65 years old; Patients did not have a history of lung diseases (Asthma, ...) or lung and heart surgeries; stable haemodynamic; Ejection fraction was more than 30 percent; Patients did not have renal failure (serum creatinine more than 2), liver disease, and history of seizure and CVA. Main exclusion criteria: Occurring unstable haemodynamic status during research; life threatening cardiac arrhythmia; Unusual hemorrhage after operation; Need to anesthesia and operation again. Sample size: 64 patients Intervention: The effect of adaptive support ventilation or synchronized mandatory intermittent ventilation on respiratory support outcomes Time of intervention: After coronary artery bypass graft surgery Outcomes: change of respiratory support outcome

#### **General information**

##### **Acronym**

##### **IRCT registration information**

IRCT registration number: **IRCT2014101919584N1**  
Registration date: **2014-12-30, 1393/10/09**  
Registration timing: **retrospective**

Last update:

Update count: **0**

##### **Registration date**

2014-12-30, 1393/10/09

##### **Registrant information**

###### **Name**

Hadi Zarei

###### **Name of organization / entity**

Isfahan University of Medical Sciences, faculty of Nursing and Midwifery

###### **Country**

Iran (Islamic Republic of)

###### **Phone**

+98 31 3442 6105

###### **Email address**

hadi.zarei767@gmail.com

##### **Recruitment status**

**Recruitment complete**

##### **Funding source**

vice chancellor for research, Isfahan university of medical sciences, Isfahan

##### **Expected recruitment start date**

2014-04-21, 1393/02/01

##### **Expected recruitment end date**

2014-06-15, 1393/03/25

##### **Actual recruitment start date**

empty

##### **Actual recruitment end date**

empty

##### **Trial completion date**

empty

##### **Scientific title**

Determining and comparing the effect of Adaptive Support Ventilation and Synchronized Intermittent Mandatory Ventilation on respiratory support outcomes

after Coronary Artery Bypass Graft surgery, Isfahan University of Medical sciences, Chamran hospital, 2014

#### Public title

Using different modes of ventilation after coronary artery bypass graft surgery

#### Purpose

Supportive

#### Inclusion/Exclusion criteria

Main inclusion criteria: Using one of the ventilation modes (Adaptive support ventilation (ASV) or synchronized intermittent mandatory ventilation (SIMV)) on Raphael ventilator was allowed by anesthesiologist; The range of age was between 25 and 65 years; Patients did not have a history of lung diseases (Asthma, ...) or lung and heart surgeries; Patients did not have renal failure (serum creatinine more than 2), liver disease, and history of seizure and CVA; left ventricular ejection fraction was more than 30 percent; In the time of ICU entrance patients were haemodynamically stable (respiratory rate less than 35 per minute, heart rate less than 150 per minute, mean atrial pressure less than 70 mmHg) and intra aortic balloon pump was not used. Main exclusion criteria: occurring unstable haemodynamic status during research (respiratory rate more than 35 per minute, heart rate more than 150 per minute, mean atrial pressure more than 70 mmHg); life threatening arrhythmia during research like ventricular tachycardia; need to excess dosages of inotropes and vasoconstrictors during research (Dopamin more than 20 milligram per hour, norepinephrin more than 0.5 milligram per hour, dobutamin more than 25 milligram per hour, and epinephrin with any doses); Unusual bleeding after operation (chest tube drainage more than 500 ml per hour, more than 350 ml per hour for 2 hours, or more than 1000 ml at all) ; Need to anesthesia and operation again.

#### Age

From **25 years** old to **65 years** old

#### Gender

Both

#### Phase

2-3

#### Groups that have been masked

*No information*

#### Sample size

Target sample size: **64**

#### Randomization (investigator's opinion)

Randomized

#### Randomization description

#### Blinding (investigator's opinion)

Single blinded

#### Blinding description

#### Placebo

Not used

#### Assignment

Parallel

#### Other design features

### Secondary Ids

empty

## Ethics committees

### 1

#### Ethics committee

##### Name of ethics committee

Moral committee of Isfahan university of medical sciences

##### Street address

Isfahan university of medical sciences, Azadi square, Hezar Jarib street, Isfahan

##### City

Isfahan

##### Postal code

8174673461

##### Approval date

2014-05-08, 1393/02/18

##### Ethics committee reference number

717/4/3

## Health conditions studied

### 1

#### Description of health condition studied

Atherosclerotic heart disease

#### ICD-10 code

I25.1

#### ICD-10 code description

Atherosclerotic heart disease

## Primary outcomes

### 1

#### Description

Intubation duration

#### Timepoint

When the intervention starts

#### Method of measurement

Hour and minutes

### 2

#### Description

Length of hospital stay

#### Timepoint

When the patient is being discharged

#### Method of measurement

Hour and minutes

## Secondary outcomes

### 1

#### Description

nothing

#### Timepoint

nothing

#### Method of measurement

nothing

## Intervention groups

### 1

#### Description

Description: In this study, patients after undergoing Coronary Artery Bypass Graft surgery (CABG) and transferring to the intensive care unit (ICU) while intubated, with approval of an anesthesiologist for possibility of using one of the Adaptive Support Ventilation (ASV) or Synchronized Intermittent Mandatory Ventilation (SIMV) modes, were randomized to the intervention or control group. Intervention group: using Adaptive Support Ventilation mode for respiratory ventilation after coronary artery bypass graft surgery in intensive care unit

#### Category

Treatment - Other

### 2

#### Description

Control group: Using Synchronized Intermittent Mandatory Ventilation mode for respiratory ventilation after coronary artery bypass graft surgery in intensive care unit

#### Category

Treatment - Drugs

## Recruitment centers

### 1

#### Recruitment center

##### Name of recruitment center

Isfahan heart diseases center

##### Full name of responsible person

Hadi Zarei - MS student of critical care nursing

##### Street address

Chamran hospital, Moshtagh 3 street, Isfahan

##### City

Isfahan

## Sponsors / Funding sources

### 1

#### Sponsor

##### Name of organization / entity

Vice chancellor for research, Isfahan university of medical sciences

##### Full name of responsible person

Dr. Peyman Adibi

##### Street address

Isfahan university of medical sciences, Hezar jarib street, Azadi square, Isfahan

##### City

Isfahan

#### 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, Isfahan 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

Isfahan university of medical sciences, faculty of nursing and midwifery

##### Full name of responsible person

Hadi Zarei

##### Position

Ms student of critical care nursing (Nurse)

##### Other areas of specialty/work

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## Person responsible for scientific inquiries

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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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**Position**  
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**Other areas of specialty/work**  
**Street address**  
Faculty of nursing and midwifery, Isfahan university of medical sciences, Hezar Jarib street, Isfahan  
**City**  
Isfahan  
**Postal code**

## 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*