

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

10 Jun 2026

### Comparison of effect of 1:1 and 1:2 frequency of Intra-aortic balloon pump on hemodynamic situation in patients with medium dose of inotrope after Coronary Artery Bypass Graft

#### Protocol summary

##### Study aim

Comparison of effect of 1:1 and 1:2 frequency of Intra-aortic balloon pump on hemodynamic situation in patients with medium dose of inotrope after Coronary Artery Bypass Graft

##### Design

quasi-experimental , with pretest - post test, no blinding,

##### Settings and conduct

Cases are patients who need Intra-aortic balloon pump after coronary artery bypass graft in Isfahan Chamran Heart Hospital, the device is set to 1: 1 frequency and it changes to 1:2 frequency for 20 minutes after reaching a median dose of inotropes, then cardiac parameters are recorded at two frequencies by echocardiography.

##### Participants/Inclusion and exclusion criteria

Entry Conditions: People who underwent CABG but due to low cardiac output, mean arterial blood pressure less than 80 mmHg failed to separate from the cardiopulmonary pump device and required postoperative IABP. Patients who are consenting to participate in the project, which will be obtained from the legal guardian if the patient is unconscious. Non-Entry Conditions: People performing CABG at the same time as aortic valve and arch-related operations such as: aortic valve replacement, Bentall surgery, etc.

##### Intervention groups

This design is a quasi-experimental study of pre-test - posttest that compares all those who were at the 1: 1 frequency before and at the start of the intervention with the 1: 2 frequency step.

##### Main outcome variables

Cardiac index; Cardiac output; Systolic blood pressure; Diastolic blood pressure; Arterial mean pressure; Heart rate; Integrated aortic blood flow at diastolic time

#### General information

##### Reason for update

##### Acronym

IABP

##### IRCT registration information

IRCT registration number: **IRCT20190922044844N1**

Registration date: **2020-03-03, 1398/12/13**

Registration timing: **registered\_while\_recruiting**

Last update: **2020-03-03, 1398/12/13**

Update count: **0**

##### Registration date

2020-03-03, 1398/12/13

##### Registrant information

##### Name

Moniresadat Afzali arani

##### Name of organization / entity

##### Country

Iran (Islamic Republic of)

##### Phone

+98 21 7753 1154

##### Email address

m.afzali@resident.mui.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2019-10-07, 1398/07/15

##### Expected recruitment end date

2020-05-20, 1399/02/31

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

### Scientific title

Comparison of effect of 1:1 and 1:2 frequency of Intra-aortic balloon pump on hemodynamic situation in patients with medium dose of inotrope after Coronary Artery Bypass Graft

### Public title

Comparison of effect of two different frequencies of Intra-aortic balloon pump on hemodynamic situation in patients after Coronary Artery Bypass Graft

### Purpose

Treatment

### Inclusion/Exclusion criteria

#### Inclusion criteria:

People who underwent CABG but due to low cardiac output, mean arterial blood pressure less than 80 mmHg failed to separate from the cardiopulmonary pump device and required postoperative IABP. Patients who are consenting to participate in the project, which will be obtained from the legal guardian if the patient is unconscious.

#### Exclusion criteria:

People who perform CABG at the same time as aortic valve and arch operations such as: aortic valve replacement, Bentall surgery, etc.

### Age

No age limit

### Gender

Both

### Phase

N/A

### Groups that have been masked

No information

### Sample size

Target sample size: **9**

More than 1 sample in each individual

Number of samples in each individual: **2**

The hemodynamic parameters defined in the article are first measured in patients with a 1: 1 frequency of the intra-aortic balloon pump and then re-measured in the phase when patients are at a 1: 2 frequency.

### Randomization (investigator's opinion)

N/A

### Randomization description

### Blinding (investigator's opinion)

Not blinded

### Blinding description

### Placebo

Not used

### Assignment

Single

### Other design features

## Secondary Ids

empty

## Ethics committees

### 1

#### Ethics committee

##### Name of ethics committee

Ethics committee of Isfahan University of Medical Sciences

##### Street address

Unit 24, No. 73, Hoquqi Ave., Somayeh Cross., Shariaty Ave.

##### City

Tehran

##### Province

Tehran

##### Postal code

1611812345

#### Approval date

2019-07-28, 1398/05/06

#### Ethics committee reference number

IR.MUI.MED.REC.1398.229

## Health conditions studied

### 1

#### Description of health condition studied

Influence of 1: 1 and 1: 2 frequency balloon pumps

#### ICD-10 code

#### ICD-10 code description

## Primary outcomes

### 1

#### Description

The rate of change in the velocity time integral of aortic blood flow during systole

#### Timepoint

In average dose of Inotropes, this parameter is measured at 1:1 frequency, then the frequency is switched to 1:2 and after 20 minutes, it's measured at 1:2 frequency again.

#### Method of measurement

The velocity time integral value of aortic blood flow during systole is measured by echocardiography.

### 2

#### Description

The measurement of area left ventricular outflow tract (area LVOT)

#### Timepoint

At moderate doses of inotropic drugs, this variable is measured before the intervention begins.

#### Method of measurement

This variable is measured by echocardiography,

### 3

#### Description

Rate of change in systolic blood pressure in patients

#### Timepoint

In average dose of Inotropes, this parameter is measured at 1:1 frequency, then the frequency is switched to 1:2

and after 20 minutes, it's measured at 1:2 frequency again.

**Method of measurement**

it's measured by a built-in arterial line for patients and cardiac monitoring devise

**4**

**Description**

Rate of change in diastolic blood pressure in patients

**Timepoint**

In average dose of Inotropes, this parameter is measured at 1:1 frequency,then the frequency is switched to 1:2 and after 20 minutes, it's measured at 1:2 frequency again.

**Method of measurement**

it's measured by a built-in arterial line for patients and cardiac monitoring devise

**5**

**Description**

Rate of change in heart rates in patients

**Timepoint**

In average dose of Inotropes, this parameter is measured at 1:1 frequency,then the frequency is switched to 1:2 and after 20 minutes, it's measured at 1:2 frequency again.

**Method of measurement**

it's measured by a cardiac monitoring devise

**6**

**Description**

Rate of change in mean arterial blood pressure in patients

**Timepoint**

In average dose of Inotropes, this parameter is measured at 1:1 frequency,then the frequency is switched to 1:2 and after 20 minutes, it's measured at 1:2 frequency again.

**Method of measurement**

it's measured by a built-in arterial line for patients and cardiac monitoring devise

**7**

**Description**

The measure of body surface area

**Timepoint**

At moderate doses of inotropic drugs, this variable is measured before the intervention begins.

**Method of measurement**

This variable is measured by echocardiography,

**8**

**Description**

Rate of change in cardiac output of patients

**Timepoint**

In average dose of Inotropes, this parameter is measured at 1:1 frequency,then the frequency is switched to 1:2 and after 20 minutes, it's measured at 1:2 frequency

again.

**Method of measurement**

It will be calculated by the formula and collected required data

**9**

**Description**

Rate of change in cardiac index of patients

**Timepoint**

In average dose of Inotropes, this parameter is measured at 1:1 frequency,then the frequency is switched to 1:2 and after 20 minutes, it's measured at 1:2 frequency again.

**Method of measurement**

It will be calculated by the formula(cardiac output divided by body surface area) and collected required data

**Secondary outcomes**

empty

**Intervention groups**

**1**

**Description**

This design is a quasi-experimental study of pre-test - posttest that compares all those who were at the 1: 1 frequency before and at the start of the intervention with the 1: 2 frequency step.

**Category**

Treatment - Devices

**Recruitment centers**

**1**

**Recruitment center**

**Name of recruitment center**

Isfahan shahid chamran hospital

**Full name of responsible person**

Amir Mirmohammadsadeghy

**Street address**

Shahid chamran Hospital, After city bridge, Third Ave, Bozorgmehr Square

**City**

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**Province**

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**Postal code**

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**Phone**

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**Email**

am\_sadeghi@med.mui.ac.ir

**Sponsors / Funding sources**

## 1

### Sponsor

**Name of organization / entity**

Esfahan University of Medical Sciences

**Full name of responsible person**

Amir Mirmohammadsadeghy

**Street address**

Isfahan University of Medical Sciences and Health Services, Hezar Jerib Ave, Azadi Square.

**City**

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**Province**

Isfahan

**Postal code**

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**Phone**

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**Email**

am\_sadeghi@med.mui.ac.ir

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

Yes

**Title of funding source**

Esfahan 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**

Esfahan University of Medical Sciences

**Full name of responsible person**

Amir mirmohammadsadeghy

**Position**

Assistant Professor of Medical School

**Latest degree**

Subspecialist

**Other areas of specialty/work**

Cardiology

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Beginning of the Tower Garden 1, Corner of the Tower, Nizwan, Isfahan

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### Person responsible for scientific inquiries

**Contact****Name of organization / entity**

Esfahan University of Medical Sciences

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**Position**

Assistant Professor of Medical School

**Latest degree**

Subspecialist

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### Person responsible for updating data

**Contact****Name of organization / entity**

Esfahan University of Medical Sciences

**Full name of responsible person**

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**Position**

Assistant Professor of Medical School

**Latest degree**

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**Email**

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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