

Clinical Trial Protocol

Iranian Registry of Clinical Trials

07 Jun 2026

The effect of erythropoietin in reduction of tissue-reperfusion injury after coronary artery baypass graft

Protocol summary

Summary

The purpose of this double blind clinical trial is to evaluate the effect of Erythropoietin on the reduction of reperfusion injury after CABG (coronary artery bypass graft). Fifty patients will be randomly assigned into two groups either to receive Erythropoietin or not. Erythropoietin will be infused 5 minutes after the end of cross clamp on surgery. Troponin and CKMB will be measured before and 10 hours after the surgery. Electrocardiography and echocardiography will be performed before and at four day and one month after the surgery.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT138809102799N1**

Registration date: **2010-01-25, 1388/11/05**

Registration timing: **registered_while_recruiting**

Last update:

Update count: **0**

Registration date

2010-01-25, 1388/11/05

Registrant information

Name

Shervin Ziabakhsh Tabary

Name of organization / entity

Mazandaran University Of Medical Sciences

Country

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Email address

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Recruitment status

Recruitment complete

Funding source

Deputy of Research and Technology of Mazandarn
University of Medical Sciences

Expected recruitment start date

2009-12-22, 1388/10/01

Expected recruitment end date

2011-01-20, 1389/10/30

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The effect of erythropoietin in reduction of tissue-reperfusion injury after coronary artery baypass graft

Public title

The effect of erythropoietin in reduction of tissue-reperfusion injury after coronary artery baypass graft

Purpose

Prevention

Inclusion/Exclusion criteria

Inclusion criteria: sign of angina pectoris need to revascularization after angiography, presence of indication for CABG Exclusion criteria: history of Myocardial Infarction since 3 months prior to the study, EF<30%, history of streptokinase taking, hemoglobin >16 , Cr > 2.5, receiving Erythropoietin since 6 months prior to the study, blood pressure > 150/95, history of Arterial fibrillation blood transfusion since 3 months prior to the study, heavy or cardiac trauma since 3 months prior to the study, polycythemia vera

Age

No age limit

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: 50

Randomization (investigator's opinion)

Randomized

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

Double blinded

Blinding description**Placebo**

Not used

Assignment

Parallel

Other design features**Secondary Ids**

empty

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

Mazandarn University of Medical Sciences

Street address

Sari

City

Sari

Postal code**Approval date**

2009-09-30, 1388/07/08

Ethics committee reference number

97-88

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

Tissue-reperfusion injury

ICD-10 code

I25.1

ICD-10 code description

Atherosclerotic heart disease

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

cardiac function (Wall Motion Score Index)

Timepoint

4 days, 30 days

Method of measurement

echocardiography

2**Description**

cardiac function (Ejection Fraction)

Timepoint

Before surgery, 4 days and 30 days after surgery

Method of measurement

echocardiography

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

cardiac marker (CKMB)

Timepoint

Before and 10 hours after surgery

Method of measurement

laboratory

2**Description**

cardiac marker (Troponin I)

Timepoint

Before and 10 hours after surgery

Method of measurement

laboratory

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

Erythropoietin (660 U/kg, 5 minutes after the end of cross clamp)

Category

Treatment - Drugs

2**Description**

no intervention in control group

Category

N/A

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

Fatemeh Alzahra Hospital

Full name of responsible person**Street address****City**

Sari

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Deputy of Research and Technology of Mazandarn
University of Medical Sciences

Full name of responsible person

Seyed Jalal Hosseinimehr

Street address

Vice-chancellor of Research and Technology of
Mazandarn University of Medical Sciences

City

Sari

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

Yes

Title of funding source

Deputy of Research and Technology of Mazandarn
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**

Mazandaran University of Medical Sciences

Full name of responsible person

Shervin Ziabakhsh Tabary

Position

Assistant professor

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Position**Other areas of specialty/work****Street address****City**

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Postal code**Phone****Fax****Email****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