

Clinical Trial Protocol

Iranian Registry of Clinical Trials

26 Jun 2026

Assessment affection of cold gel pack application on the sternal incision and pleural chest tube insertion site on the pulmonary function and pain due to coughing and deep breathing in coronary artery bypass graft surgery patients

Protocol summary

Summary

Study Objective: Assessment affection of cold gel pack application on the sternal incision and pleural chest tube insertion site on the pulmonary function and pain due to coughing and deep breathing in coronary artery bypass graft surgery patients The trial will be conducted randomized parallel design Study Population: Patients undergoing open heart surgery in Madani hospital A) Inclusion Criteria 1- The patient consent to participate in the study. 2- The patient is over 40 years old. 3- The patient is able to read and write. 4- The patient can speak Farsi and Azari. 5- The patient orient to person, place and time. 6- The patient is able t use pain scale. 7- The patient has been underwent coronary artery bypass surgery (with or without valve replacement) with median sternotomy located. B) Exclusion criteria 1- The patient has a history of allergy or sensitivity to cold. 2- The patient has a history of Raynaud 's syndrome , sickle cell anemia , Cryoglobulinemia , diabetes and delirium. 3- The patient has severe visual, hearing and speech impairment. 4- The patient is connected to a ventilator at the time of data collection. 5- The patient has been ill at the time of data collection. 6- The patient has postoperative complications such as infection , bleeding , uncontrolled atrial fibrillation and opening the wound edges. The sample size: 25 people for each groups. In this study, patients who underwent coronary artery bypass surgery with median sternotomy randomly used cooling gel bag or gel bag with room temperature. Data on the afternoon of the first day of practice will be collected. The pain in both groups will be measured before of use of the gel pack and at the end coughing and deep breathing by the researcher with numerical pain rating scale. lung volumes are measured by Spirometry in three times before surgery and before and after applying the gel and coughing and deep. study

time: 20 minutes. Expected Outcome: change in pain and pulmonary function results.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT201502049422N9**

Registration date: **2015-03-15, 1393/12/24**

Registration timing: **prospective**

Last update:

Update count: **0**

Registration date

2015-03-15, 1393/12/24

Registrant information

Name

Aram Feizi

Name of organization / entity

Urmia University of Medical Sciences

Country

Iran (Islamic Republic of)

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+98 44 1336 5836

Email address

aramfeizi@umsu.ac.ir

Recruitment status

Recruitment complete

Funding source

Urmia Medical Science University

Expected recruitment start date

2015-03-21, 1394/01/01

Expected recruitment end date

2015-05-22, 1394/03/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Assessment affection of cold gel pack application on the sternal incision and pleural chest tube insertion site on the pulmonary function and pain due to coughing and deep breathing in coronary artery bypass graft surgery patients

Public title

Effects of cold on the pain

Purpose

Treatment

Inclusion/Exclusion criteria

A) Inclusion Criteria 1- The patient consent to participate in the study. 2- The patient is over 40 years old. 3- The patient is able to read and write. 4- The patient can speak Farsi and Turkish. 5- The patient orient to person, place and time. 6- The patient is able t use pain scale. 7- The patient has been underwent coronary artery bypass surgery (with or without valve replacement) with median sternotomy located. B) Exclusion criteria 1- The patient has a history of allergy or sensitivity to cold. 2- The patient has a history of Raynaud 's syndrome , sickle cell anemia , Cryoglobulinemia , diabetes and delirium. 3- The patient has severe visual, hearing and speech impairment. 4- The patient is connected to a ventilator at the time of data collection. 5- The patient has been ill at the time of data collection. 6- The patient has postoperative complications such as infection , bleeding , uncontrolled atrial fibrillation and opening the wound edges.

Age

From **40 years** old to **100 years** old

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)**

Single blinded

Blinding description**Placebo**

Used

Assignment

Parallel

Other design features**Secondary Ids**

empty

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

Tabriz Medical Science University

Street address

Central Building of Medical science, Golgasht Avenue, Azadi Street, Tabriz, East Azarbaijan

City

Tabriz

Postal code**Approval date**

2015-01-26, 1393/11/06

Ethics committee reference number

93160

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

Pain in patients undergoing coronary artery surgery

ICD-10 code

R07.1

ICD-10 code description

Chest pain on breathing

2**Description of health condition studied**

pulmonary function tests

ICD-10 code

R94.2

ICD-10 code description

Abnormal results of pulmonary function studies

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

pain

Timepoint

before and 20 minutes after intervention

Method of measurement

numeric pain scale

2**Description**

pulmonary function results

Timepoint

before of operation /before of use of ice packed/ 20 minutes after use of ice packed

Method of measurement

spirometry

Secondary outcomes

empty

Intervention groups

1

Description

In this study, the cooling gel pack with dimensions (10 x 25 cm) that is placed at room temperature and the timer is set for 20 minutes for the control group. Then, gel pack is placed in the dressing on the sternum incision site and around the pleural chest tube. A pillow or folded sheet will be given to the patient to support and immobilize the wound site before coughing or deep breathing. Gel pack is removed in 20 minutes and head of the bed elevated to 45-90 degrees for coughing and deep breathing. Pain intensity will be measured by a researcher using numerical pain scale, before and after applying gel pack at the end of coughing and deep breathing in the control group. The lung volumes are measured by using spirometry three times; before surgery, before and after applying the gel pack and coughing and deep breathing. Data will be collected in the afternoon of the first day after the operation.

Category

Treatment - Other

2

Description

In this study, the cooling gel pack with dimensions (10 x 25 cm) that has been frozen for at least one hour (reaching a pack's temperature between 0 to - 5°C) is removed from the freezer and the timer is set for 20 minutes in the intervention group. Then, gel pack is placed in the dressing on the sternum incision site and around the pleural chest tube. A pillow or folded sheet will be given to the patient to support and immobilize the wound site before coughing or deep breathing. Gel pack is removed in 20 minutes and head of the bed elevated to 45-90 degrees for coughing and deep breathing. Pain intensity will be measured by a researcher using numerical pain scale, before and after applying the gel pack at the end of coughing and deep breathing in the intervention group. The lung volumes are measured by using spirometry three times; before surgery, before and after applying the gel pack, and coughing and deep breathing. Data will be collected in the afternoon of the first day after the operation

Category

Treatment - Other

Recruitment centers

1

Recruitment center

Name of recruitment center

Madani hospital

Full name of responsible person

Feizi Aram

Street address

Madani hospital, In front of Research and Development complex of University, University

Street, Tabriz, East Azarbaijan

City

Tabriz

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Urima Medical Science University

Full name of responsible person

Mohebi Iraj

Street address

Research and Technology Assistance Building, University Staff, End of The Emergency Center, Resalat Blvd., Urmia, West Azarbaijan

City

Urmia

Grant name

Grant code / Reference number

Is the source of funding the same sponsor organization/entity?

Yes

Title of funding source

Urima Medical Science University

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

Madani hospital, Tabriz Medical Science University

Full name of responsible person

Noorellahi Davod

Position

Msc in Critical Care Nursing

Other areas of specialty/work

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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