

Clinical Trial Protocol

Iranian Registry of Clinical Trials

21 Jul 2026

The Effect of Desmopressin on The Post Operative Bleeding in Patients Undergoing Open Heart Surgery

Protocol summary

Study aim

Reduced postoperative bleeding in open heart surgery patients

Design

Clinical trial with control group, with parallel groups, double-blind, randomized, phase 2 on 194 patients.

Settings and conduct

All patients undergoing open heart surgery will be included in the study, according to a specialist in cardiovascular surgery. An expert who will be responsible for measuring bleeding will be blinded in terms of knowledge of the type of medication. Desmopressin and Sodium Chloride 0.9 sprays are given by the anesthesiologist in the operating room 30 minutes before the operation at a rate of 0.3 micrograms per kilogram, respectively, to each patient in the intervention and control group. The person giving the medicine in the operating room and the person who measures the patient's bleeding in the ICU OH section are the same for both groups and do not know what group the patient is in and what medication he has received

Participants/Inclusion and exclusion criteria

All patients are candidates for open heart surgery subject to informed consent Conscious dissatisfaction

Intervention groups

The intervention group will receive 0.3 micrograms per kilogram of desmopressin half an hour before the start of surgery. The control group will receive 0.3 micrograms per kilogram of 0.9 sodium chloride half an hour before the start of surgery.

Main outcome variables

Postoperative bleeding

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20200424047185N1**

Registration date: **2020-05-26, 1399/03/06**

Registration timing: **registered_while_recruiting**

Last update: **2020-05-26, 1399/03/06**

Update count: **0**

Registration date

2020-05-26, 1399/03/06

Registrant information

Name

Hamed Kiani

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 54 3328 4340

Email address

kianihamed93@yahoo.com

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2020-03-20, 1399/01/01

Expected recruitment end date

2021-03-19, 1399/12/29

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The Effect of Desmopressin on The Post Operative Bleeding in Patients Undergoing Open Heart Surgery

Public title

The Effect of Desmopressin on The Post Operative Bleeding

Purpose

Prevention

Inclusion/Exclusion criteria**Inclusion criteria:**

Candidate for open heart surgery

Exclusion criteria:

Blood coagulation disease

Age

No age limit

Gender

Both

Phase

3

Groups that have been masked

- Care provider
- Outcome assessor

Sample size

Target sample size: **194**

Randomization (investigator's opinion)

Randomized

Randomization description

Easy (available) sampling, in which two groups of age and sex will be matched and matched frequently using stratified randomization. The explanation of this method is that after entering the study, patients are divided into two groups, male and female, which in each group of patients are divided into two categories of patients under 50 years of age and over 50 years of age. Depending on the accident, individuals will be assigned to two intervention groups and a control group.

Blinding (investigator's opinion)

Double blinded

Blinding description

An expert who will be responsible for measuring bleeding reduction will be blinded in terms of knowledge of the type of medication. Desmopressin and normal saline sprays are given to each of the patients in the intervention or control group 30 minutes before the operation, respectively, by the anesthesiologist in the operating room 30 minutes before the start of the operation. The pharmacist and the person measuring the patient's bleeding at ICUOH are the same for both groups and do not know what group the patient is in and what medication he or she has received (they are blind).

Placebo

Used

Assignment

Parallel

Other design features**Secondary Ids**

empty

Ethics committees**1****Ethics committee**

Name of ethics committee

Ethics Committee of Zahedan University of Medical Sciences

Street address

Dr. Hesabi Square, Medical Sciences Campus

City

Zahedan

Province

Sistan-va-Balouchestan

Postal code

9816743463

Approval date

2020-04-21, 1399/02/02

Ethics committee reference number

IR.ZAUMS.REC.1399.027

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

Bleeding - open heart surgery - Desmopressin

ICD-10 code**ICD-10 code description****Primary outcomes****1****Description**

Postoperative bleeding

Timepoint

Measure the patient's bleeding rate two, six, eight, twelve, and twenty-four hours after surgery

Method of measurement

The amount of blood in the Chest Bottle is in milliliters

Secondary outcomes

empty

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

Intervention group: Desmopressin is an artificial analogue of vasopressin group. 0.3 micrograms per kilogram is given to each patient of the intervention group 30 minutes before the operation.

Category

Treatment - Surgery

2**Description**

Control group: Sodium chloride solution is 0.9% injectable, a sterile, non-febrile solution, free of preservatives and isotonic, and contains 9 mg of sodium chloride per ml. 0.3 micrograms per kilogram is given to each patient in the control group.

Category

Prevention

Recruitment centers

1

Recruitment center

Name of recruitment center

Ali Ibn Abi Talib Hospital

Full name of responsible person

Azadi Ahmad Abadi Changiz

Street address

Salamat Blvd - Persian Gulf Highway

City

Zahedan

Province

Sistan-va-Balouchestan

Postal code

9816743111

Phone

+98 54 3329 5570

Email

public@zaums.ac.ir

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Zahedan University of Medical Sciences

Full name of responsible person

AzadiAhmad Abadi Changiz

Street address

Campus of Medical Sciences - Dr. Hesabi Square

City

Zahedan

Province

Sistan-va-Balouchestan

Postal code

9816743463

Phone

+98 54 3329 5744

Email

public@zaums.ac.ir

Grant name

Grant code / Reference number

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

Yes

Title of funding source

Zahedan University of Medical Sciences

Proportion provided by this source

100

Public or private sector

Private

Domestic or foreign origin

Domestic

Category of foreign source of funding

empty

Country of origin

Type of organization providing the funding

Persons

Person responsible for general inquiries

Contact

Name of organization / entity

Zahedan University of Medical Sciences

Full name of responsible person

Azadiyahmadabadi Changiz

Position

Assistant Professor

Latest degree

Subspecialist

Other areas of specialty/work

Heart surgery

Street address

Salamat Blvd-Persian Gulf Highway

City

Zahedan

Province

Sistan-va-Balouchestan

Postal code

9816743111

Phone

+98 54 3329 5570

Email

kianihamed93@yahoo.com

Person responsible for scientific inquiries

Contact

Name of organization / entity

Zahedan University of Medical Sciences

Full name of responsible person

Azadiyahmadabadi Changiz

Position

Assistant Professor

Latest degree

Subspecialist

Other areas of specialty/work

Heart surgery

Street address

Salamat Blvd-Persian Gulf Highway

City

Zahedan

Province

Sistan-va-Balouchestan

Postal code

9816743111

Phone

+98 54 3329 5570

Email

kianihamed93@yahoo.com

Person responsible for updating data

Contact

Name of organization / entity

Zahedan University of Medical Sciences

Full name of responsible person

Azadiyahmadabadi Changiz

Position

Assistant Professor

Latest degree

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Other areas of specialty/work

Heart surgery

Street address

Salamat Blvd-Persian Gulf Highway

City

Zahedan

Province

Sistan-va-Balouchestan

Postal code

9816743111

Phone

0098 54 3329557

Email

kianihamed93@yahoo.com

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

Sharing plan**Deidentified Individual Participant Data Set (IPD)**