

Clinical Trial Protocol

Iranian Registry of Clinical Trials

13 Jun 2026

The Effect of Aminophylline on Renal Function after Cardiac Surgery

Protocol summary

Summary

Objectives: the aim of this study was to assess the effectiveness of aminophylline for prevention of renal impairment after cardiac surgery. Inclusion criteria: all patients undergoing cardiac surgery and can use Aminophylline. Research Design: Conduction of a randomized, double-blind, clinical trial, method of randomization: simple. Exclusion criteria: patients with cardiac arrhythmia, tachycardia, and patients with kidney damage in case of non- oliguric, daily urine of patients who have obstructive uropathy, patients with congenital renal anomalies. One hundred and fourthly-four patients undergoing cardiac surgery were risk stratified as per Cleveland score to assess for prediction of AKI(acute kidney injury). 72 patients in the intervention group received a bolus aminophylline of 5 mg/kg and a subsequent continuous infusion of 0.25 mg/kg/h for up to 48 h, while, 72 Patients of control group just received usual postoperative care. Serum creatinine concentrations were measured preoperatively and daily until 2 days after surgery and the glomerular filtration rate estimated using Cockcroft and Gault formula. Hourly urine output was recorded and patients assigned to respective RIFLE stage of AKI. Cleveland score ≥ 6 was associated with higher incidence of AKI: I and F. Number needed to treat, an insight into the clinical relevance of a specific treatment, is 8.

General information

Acronym

acute kidney injury (AKI), acute renal failure (ARF)

IRCT registration information

IRCT registration number: **IRCT2015010619470N10**

Registration date: **2015-01-27, 1393/11/07**

Registration timing: **retrospective**

Last update:

Update count: **0**

Registration date

2015-01-27, 1393/11/07

Registrant information

Name

Farzaneh Masihi

Name of organization / entity

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Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Vice Chancellor for Research, Shiraz University of Medical

Expected recruitment start date

2014-10-23, 1393/08/01

Expected recruitment end date

2015-01-21, 1393/11/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The Effect of Aminophylline on Renal Function after Cardiac Surgery

Public title

The Effect of Aminophylline on Renal after Cardiac Surgery

Purpose

Prevention

Inclusion/Exclusion criteria

Inclusion criteria: all patients undergoing cardiac surgery and can use Aminophylline. Exclusion criteria: patients with cardiac arrhythmia, tachycardia, patients with kidney damage in case of non oliguric, daily urine of

patients who have obstructive uropathy, patients with congenital renal anomalies

Age

No age limit

Gender

Both

Phase

2

Groups that have been masked

No information

Sample size

Target sample size: **144**

Randomization (investigator's opinion)

Randomized

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

Double blinded

Blinding description**Placebo**

Used

Assignment

Parallel

Other design features**Secondary Ids**

empty

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

Ethic Committee of Shiraz University Of Medical Sciences

Street address

Central Building of Shiraz University of Medical Sciences, Zand Street

City

Shiraz

Postal code**Approval date**

2010-08-20, 1389/05/29

Ethics committee reference number

CP-P-9374-7224

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

Cardiac Surgeries

ICD-10 code

197.1

ICD-10 code description

Cardiac insufficiency Heart failure following cardiac surgery or due to presence of cardiac prosthesis

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

Diagnosis of Kidney Damage

Timepoint

Daily measurement: On the days before and after surgery and the second and third days after surgery.

Method of measurement

RIFLE scale: It based on the amount of creatinine and GFR calculated by the amount of kidney damage.

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

Diagnosis of Kidney Damage

Timepoint

Daily measurement: On the days before and after surgery and the second and third days after surgery

Method of measurement

Urine Output

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

Patients in the intervention group received a bolus aminophylline of 5 mg/kg before the surgery and a subsequent continuous infusion of 0.25 mg/kg/h for up to 48 h after the surgery.

Category

Treatment - Drugs

2**Description**

Control group: patients in the control group didn't take aminophylline.

Category

Placebo

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

Kowsar Hospital

Full name of responsible person

Peyman Alishahi

Street address

Ghasrodasht Street

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Vice Chancellor for Research, Shiraz University of Medical Sciences

Full name of responsible person

Dr Seed Basir Hashemi

Street address

Vice Chancellor for Research, Shiraz University of Medical Sciences, 7th floor, Central Building of Shiraz University of Medical Sciences, Zand Street

City

Shiraz

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, Shiraz 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**

Shiraz University of Medical Sciences

Full name of responsible person

Peyman Alishahi

Position

Physician, Anesthesiology Resident

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Person responsible for updating data

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Full name of responsible person

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Position

English Consultant, Anesthesiology Research Center Staff

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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