

Clinical Trial Protocol

Iranian Registry of Clinical Trials

09 Jun 2026

The effect of curcumin in the prevention of contrast induced nephropathy in patients with moderate kidney failure following coronary angioplasty or angiography

Protocol summary

Summary

Contrast nephropathy is the most common cause of iatrogenic acute kidney injury. This problem particularly in CKD patients is more common. In various studies, the positive effect of antioxidants in preventing of it is suggested. Therefore, we decided to study the antioxidant effect of curcumin in the prevention of contrast nephropathy in moderate chronic kidney disease patients. All patients with stable angina and unstable or non-ST elevated MI in the of Ghaem and Imam Reza hospital who are candidates for elective and/or emergency coronary angiography or angioplasty enrolled in study, if creatinine more than 1.2 or GFR between 30 to 60 ml / min are based on prior lab tests. It is done as a double blind study for both patients and physicians. 30 patients will be randomly allocated to receive capsule Curcumin 500 mg three times daily from 2 days before angiography until 3 days after it and 30 patients in control group will receive placebo to the same extent. In both groups, from 12 hours before angiography normal saline solution infuse at a rate of 1ml/Kg/hr and it will continue to 12 hours after that. Serum creatinine will be reevaluate in second and third days after angiography. Urinary NGAL will be assessed in the next day of angiography.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT138706261256N1**

Registration date: **2017-10-24, 1396/08/02**

Registration timing: **retrospective**

Last update:

Update count: **0**

Registration date

2017-10-24, 1396/08/02

Registrant information

Name

Maryam Hami

Name of organization / entity

Health Organization

Country

Iran (Islamic Republic of)

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Email address

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Recruitment status

Recruitment complete

Funding source

Mashhad University of Medical Sciences

Expected recruitment start date

2016-10-11, 1395/07/20

Expected recruitment end date

2017-05-01, 1396/02/11

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The effect of curcumin in the prevention of contrast induced nephropathy in patients with moderate kidney failure following coronary angioplasty or angiography

Public title

The effect of curcumin in the prevention of kidney failure following heart angiography

Purpose

Prevention

Inclusion/Exclusion criteria

Inclusion criteria: All patients undergoing elective coronary angiography who are older than 18 years and are suffering from chronic kidney failure. Exclusion criteria: patients on dialysis, acute renal failure before angiography, cardiogenic shock, heart failure, use of medications such as N-acetyl cysteine, vitamin C, use of intravenous contrast to other causes, mannitol, diuretics, theophylline, warfarin and dopamine in the past two weeks.

Age

From **18 years** old

Gender

Both

Phase

2-3

Groups that have been masked

No information

Sample size

Target sample size: **60**

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

Mashhad University Of Medical sciences ethics Committee

Street address

Vice President of Research, Ghoreshi building, Daneshgah st.

City

Mashhad

Postal code

Approval date

2016-08-17, 1395/05/27

Ethics committee reference number

IR.MUMS.sm.REC.1395.220

Health conditions studied

1

Description of health condition studied

Acute kidney injury

ICD-10 code

N17, N18,

ICD-10 code description

Renal Failure

Primary outcomes

1

Description

serum creatinine

Timepoint

0,1,3 days- 1th day

Method of measurement

mg/dl

2

Description

urinary NGAL

Timepoint

First day

Method of measurement

ng/ml

Secondary outcomes

1

Description

nausea

Timepoint

daily

Method of measurement

history & physical exam

2

Description

diarrhea

Timepoint

daily

Method of measurement

history & physical exam

Intervention groups

1

Description

Control group receive placebo capsule 500 mg three times daily from 2 days before angiography until 3 days after it.

Category

Treatment - Drugs

2

Description

Intervention group receive Curcumin capsule 500 mg three times daily from 2 days before angiography until 3 days after it.

Category

Treatment - Drugs

Recruitment centers1**Recruitment center****Name of recruitment center**

Ghaem Hospital

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

Mashhad

2**Recruitment center****Name of recruitment center**

Emam Reza Hospital

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

Mashhad

Sponsors / Funding sources1**Sponsor****Name of organization / entity**Vice Chancellor Of Research, Mashad University of
Medical Sciences**Full name of responsible person**

Mohsen Tafaghodi

Street addressVice President of Research- Ghoreshi building-
Daneshgah st.**City**

Mashhad

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

Yes

Title of funding sourceVice Chancellor Of Research, Mashad 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**Vice Chancellor Of Research, Mashad University of
Medical Sciences**Full name of responsible person**

Mohsen Tafaghodi

Position

Vice Chancellor Of Research

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Web page address**Person responsible for scientific
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Maryam Hami

Position

Associated Professor Of Nephrology

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Full name of responsible person

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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