

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

09 Jul 2026

### Effect of N - acetylcysteine in the prevention of ototoxicity caused by vancomycin in the treatment of peritonitis in patients undergoing peritoneal dialysis.

#### Protocol summary

##### Summary

The aim of this study is to investigate the possible protective effects of N - acetylcysteine on the incidence of hearing damage caused by vancomycin in the treatment of peritonitis in patients undergoing peritoneal dialysis. Inclusion criteria are the patients who undergo peritoneal dialysis and have peritonitis. Exclusion criteria is the patients who have hearing damage before our study. After informed consent is obtained , 60 patients under peritoneal dialysis with peritonitis are enrolled. All patients treated by I.P. vancomycin within ISPD protocol. Patients were divided into two groups : Groups recving N - acetylcysteine (30 patients) and those who do not take N - acetylcysteine (30 patients). Patients undergoing audiometric studies at frequencies of 250 , 500 , 1000 , 2000 , 3000 , 4000 , 6000 and 8000. The control group treated with vancomycin and other group treated with I.P. vancomycin and N - acetylcystein orally 600 mg BID. After 21 day treatment period , two groups undergoing audiometric studies and hearing impairment will be assessed.

#### General information

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT201202148032N2**  
Registration date: **2013-07-09, 1392/04/18**  
Registration timing: **registered\_while\_recruiting**

Last update:

Update count: **0**

##### Registration date

2013-07-09, 1392/04/18

##### Registrant information

##### Name

Sima Abediazar

##### Name of organization / entity

Tabriz University Of Medical Sciences

##### Country

Iran (Islamic Republic of)

##### Phone

+98 41 1334 7054

##### Email address

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##### Recruitment status

**Recruitment complete**

##### Funding source

Tabriz University of Medical Sciences

##### Expected recruitment start date

2012-09-28, 1391/07/07

##### Expected recruitment end date

2013-11-28, 1392/09/07

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

##### Scientific title

Effect of N - acetylcysteine in the prevention of ototoxicity caused by vancomycin in the treatment of peritonitis in patients undergoing peritoneal dialysis.

##### Public title

Effect of N - Acetylcysteine in the prevention of ototoxicity caused by vancomycin.

##### Purpose

Prevention

##### Inclusion/Exclusion criteria

Inclusion criteria : Patients with peritoneal dialysis because of kidney problems ; Patients who do not use

ototoxic drugs ; Patients who have no previous hearing impairment Exclusion criteria : Baseline audiometry with hearing damage ; Patients who use other ototoxic drugs

#### Age

No age limit

#### Gender

Both

#### Phase

N/A

#### Groups that have been masked

No information

#### Sample size

Target sample size: 60

#### Randomization (investigator's opinion)

Randomized

#### Randomization description

#### Blinding (investigator's opinion)

Single blinded

#### Blinding description

#### Placebo

Not used

#### Assignment

Parallel

#### Other design features

## Secondary Ids

empty

## Ethics committees

### 1

#### Ethics committee

##### Name of ethics committee

Tabriz University of Medical Sciences

##### Street address

Tabriz - University Ave. - Tabriz University of Medical Sciences

##### City

Tabriz

##### Postal code

##### Approval date

2012-12-17, 1391/09/27

##### Ethics committee reference number

91194

## Health conditions studied

### 1

#### Description of health condition studied

Peritonitis

#### ICD-10 code

K65.9

#### ICD-10 code description

Peritonitis, unspecified

## Primary outcomes

### 1

#### Description

Autotoxicity

#### Timepoint

Baseline and 21 days after treatment with N-acetylcysteine

#### Method of measurement

Audiometry

## Secondary outcomes

empty

## Intervention groups

### 1

#### Description

Intervention: Receiving vancomycin intraperitoneal 1g q 5 days plus N - acetylcysteine 600 mg BID orally for 21 days.

#### Category

Treatment - Drugs

### 2

#### Description

Control: Receiving vancomycin intraperitoneal 1g q 5 days for 21 days.

#### Category

Treatment - Drugs

## Recruitment centers

### 1

#### Recruitment center

##### Name of recruitment center

Clinic Of Nephrology, Imam Reza Hospital

##### Full name of responsible person

Sima Abediazar

##### Street address

Imam Reza Hospital, Golgasht Ave.

##### City

Tabriz

## Sponsors / Funding sources

### 1

#### Sponsor

##### Name of organization / entity

Research Assistance Of Tabriz University Of Medical Science

##### Full name of responsible person

Mohammad Reza Rashidi

##### Street address

Research Assistance, Tabriz University Of Medical Science, Golgasht Ave.

##### City

Tabriz

**Grant name**  
**Grant code / Reference number**  
**Is the source of funding the same sponsor organization/entity?**  
Yes  
**Title of funding source**  
Research Assistance Of Tabriz University Of Medical Science  
**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

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## Person responsible for updating data

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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*