

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

10 Jun 2026

### Pain Relieving Effect of Glycerol Trinitrate (GTN) in Renal Colic: a Randomized Placebo-Controlled Trial

#### Protocol summary

##### Study aim

Evaluation of pain relieving effect of GTN Glyceryl Trinitrate spray in patients referred to the emergency department (ED) due to renal colic pain

##### Design

This study will be a blind, randomized, placebo-controlled three-way trial.

##### Settings and conduct

This study will be performed in the internal ward of Shariati Hospital, Tehran, Iran. Patients entered the study after signing the informed consent form. Researchers will adhere to the Helsinki Declaration of Principles during the study.

##### Participants/Inclusion and exclusion criteria

Patients over 16 years of age, who are clinically diagnosed by an emergency physician with imaging (ultrasound or CT scan) to have renal colic and have a pain level higher than 5 on the VAS scale). The following will be considered as exclusion criteria: symptoms of peritoneal irritation, pregnancy and lactation, fever or hypotension; History of sildenafil, tadalafil, vardenafil, antihypertensive drugs, aspirin, antimuscarinic, alcohol, ergotamine, haloperidol or phenothiazine during the last 4 weeks,

##### Intervention groups

The intervention group receiving the GTN Glyceryl Trinitrate spray and the control group that will receive only the placebo treatment.

##### Main outcome variables

Pain relief is defined as the reduction of VAS pain to less than 5. Therefore, the pain will be measured again in patients after 5 minutes, and if the VAS pain was still more than 5, the patient will receive another 30 mg of ketorolac injection. Pain will be re-estimated 30 minutes after this dose, and if the patient has a VAS pain score higher than 5, the next dose of ketorolac will be injected. Response to VAS treatment is defined as less than 5.

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20180428039443N2**

Registration date: **2021-04-06, 1400/01/17**

Registration timing: **registered\_while\_recruiting**

Last update: **2021-04-06, 1400/01/17**

Update count: **0**

##### Registration date

2021-04-06, 1400/01/17

##### Registrant information

##### Name

Ahmad Reza Dehpour

##### Name of organization / entity

##### Country

Iran (Islamic Republic of)

##### Phone

+98 21 8897 3654

##### Email address

dehpour@yahoo.com

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2021-02-12, 1399/11/24

##### Expected recruitment end date

2021-09-20, 1400/06/29

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

##### Scientific title

Pain Relieving Effect of Glycerol Trinitrate (GTN) in Renal Colic: a Randomized Placebo-Controlled Trial

#### Public title

Pain Relieving Effect of Glycerol Trinitrate (GTN) in Renal Colic

#### Purpose

Treatment

#### Inclusion/Exclusion criteria

##### Inclusion criteria:

Patients over 16 years of age, who are clinically diagnosed by an emergency physician with imaging (ultrasound or CT scan) to have renal colic and have a pain level higher than 5 on the VAS scale). Patients who have treated themselves or had side effects from diseases such as gastrointestinal, heart, kidney or liver are not included. The following will be considered as exclusion criteria: symptoms of peritoneal irritation, pregnancy and lactation, fever or hypotension; History of sildenafil, tadalafil, vardenafil, antihypertensive drugs, aspirin, antimuscarinic, alcohol, ergotamine, haloperidol or phenothiazine during the last 4 weeks,

##### Exclusion criteria:

allergic reaction to GTN; Severe clinical anemia; Recent concussion or any damage to the central nervous system, malnutrition, hypothyroidism; Hypothermia; And drug abusers.

#### Age

From **16 years** old

#### Gender

Both

#### Phase

N/A

#### Groups that have been masked

- Participant
- Care provider
- Data analyser

#### Sample size

Target sample size: **100**

#### Randomization (investigator's opinion)

Randomized

#### Randomization description

The random number sequences of volunteers are made by the Sealed Envelope | Randomization site. Using random quadrilateral blocks, the random chain is created at first by the number of sample volumes.

#### Blinding (investigator's opinion)

Triple blinded

#### Blinding description

Because of the nature of the outcome of the disease (pain), blindness or blindness does not disclose the study. However, the final data for analysis will be coded and analyzed without the knowledge of the treatment group. Data collection authorities and those who evaluate the outcome, Data Safety and Monitoring Committee, and those who prepare the draft article, are kept blind to the studied groups.

#### Placebo

Used

#### Assignment

Parallel

#### Other design features

This study will be a blind, randomized, placebo-controlled three-way trial that will be performed in the internal ward of Shariati Hospital, Tehran, Iran. Patients entered the study after signing the informed consent form.

Researchers will adhere to the Helsinki Declaration of Principles during the study.

#### Secondary Ids

empty

#### Ethics committees

##### 1

##### Ethics committee

###### Name of ethics committee

Tehran University of Medical Sciences

###### Street address

Medical School, PourSina St., Tehran, 1417613151, Iran

###### City

Tehran

###### Province

Tehran

###### Postal code

1417613151

##### Approval date

2021-01-29, 1399/11/10

##### Ethics committee reference number

IR.TUMS.MEDICINE.REC.1399.1034

#### Health conditions studied

##### 1

##### Description of health condition studied

Renal Colic

##### ICD-10 code

N23

##### ICD-10 code description

Unspecified renal colic

#### Primary outcomes

##### 1

##### Description

Pain Relieving

##### Timepoint

Pain will be measured again in patients after 5 minutes, and if the VAS pain is still more than 5, the patient will receive another 30 mg of ketorolac injection. Pain will be re-estimated 30 minutes after this dose, and if the patient has a VAS pain score higher than 5, the next dose of ketorolac will be injected.

##### Method of measurement

Response to VAS treatment is defined as less than 5.

## Secondary outcomes

empty

## Intervention groups

### 1

#### Description

Intervention group: Patients who receive, in addition to standard treatment Glycerol Trinitrate (GTN). GTN sprays contain 0.4 mg of trinitroglycerin.

#### Category

Treatment - Drugs

### 2

#### Description

Control group: Patients undergoing standard treatment and receiving placebo. Placebo sprays will contain exponents to have a similar odor to the patients receiving them until they look exactly the same. We will record the number for each package and content as a placebo or GTN spray.

#### Category

Placebo

## Recruitment centers

### 1

#### Recruitment center

##### Name of recruitment center

Shariati Hospital

##### Full name of responsible person

Javad Seyedhosseini

##### Street address

Medical School, PourSina St., Tehran, 1417613151, Iran

##### City

Tehran

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##### Postal code

1417613151

##### Phone

+98 21 8987 3652

##### Email

jshosseini@gmail.com

##### Web page address

<https://scholar.google.com/citations?user=40HkrBcAAAJ&hl=en>

## Sponsors / Funding sources

### 1

#### Sponsor

##### Name of organization / entity

Tehran University of Medical Sciences

##### Full name of responsible person

Amir Esmaeil Saghafi nia

#### Street address

Medical School, PourSina St., Tehran, 1417613151, Iran

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#### Email

saghafinia@dorsadarou.com

#### Grant name

#### Grant code / Reference number

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

No

#### Title of funding source

Tehran 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

Industry

## Person responsible for general inquiries

#### Contact

##### Name of organization / entity

Tehran University of Medical Sciences

##### Full name of responsible person

Ahmad Reza Dehpour

##### Position

Professor

##### Latest degree

Ph.D.

##### Other areas of specialty/work

Medical Pharmacy

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##### Email

dehpour@yahoo.com

## Person responsible for scientific

## **inquiries**

### **Contact**

**Name of organization / entity**

Tehran University of Medical Sciences

**Full name of responsible person**

Ahmad Reza Dehpour

**Position**

Professor

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**Other areas of specialty/work**

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## **Person responsible for updating data**

### **Contact**

**Name of organization / entity**

Tehran University of Medical Sciences

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Razieh Mohammad Jafari

**Position**

Assistant Professor

**Latest degree**

Ph.D.

**Other areas of specialty/work**

Medical Pharmacy

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rmjafari@sina.tums.ac.ir

## **Sharing plan**

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

Yes - There is a plan to make this available

**Study Protocol**

Yes - There is a plan to make this available

**Statistical Analysis Plan**

Yes - There is a plan to make this available

**Informed Consent Form**

Yes - There is a plan to make this available

**Clinical Study Report**

Yes - There is a plan to make this available

**Analytic Code**

Yes - There is a plan to make this available

**Data Dictionary**

Yes - There is a plan to make this available

**Title and more details about the data/document**

All patient information documented data without identification. The information entered will eventually be entered in the database of the database.

**When the data will become available and for how long**

Data will be available upon national and international patenting process completion of the study.

**To whom data/document is available**

All interested researchers

**Under which criteria data/document could be used**

There is no specific condition.

**From where data/document is obtainable**

Dr. Ahmad Reza Dehpour

**What processes are involved for a request to access data/document**

The procedure outlined by the Ethics Committee of Tehran University of Medical Sciences

**Comments**