

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

11 Jun 2026

### Role of thiamine in the management of post-operative analgesia in maxillofacial surgery.

#### Protocol summary

##### Summary

(1) Objectives: Thiamine is a water-soluble vitamin of the B complex. First named aneurin for the negative neurological effects if not present in the diet, it was finally assigned the generic name vitamin B1. Thiamine plays a key role in certain metabolic reaction, including: nervous system and muscle functioning; flow of electrolytes in and out of nerve and muscle cells (through ion channels); multiple enzyme processes (via the coenzyme thiamin pyrophosphate); carbohydrate metabolism; and production of hydrochloric acid (which is necessary for proper digestion). Thiamine is generally considered safe and relatively nontoxic, even at high doses. Thiamine also helps convert carbohydrate to fat for storage of potential energy. Thiamine is a major factor in the metabolism of carbohydrate, fat, amino acid, glucose, and alcohol . It is a co-factor of pyruvate dehydrogenase and necessary for aerobic metabolism , that may increase oxygen extraction in critically ill patients. The purpose of this study was to evaluate the ability of intravenous thiamine for postoperative pain reduction in oral and maxillofacial surgery. (2) Design: This is a double-blind randomised clinical trial study, that evaluate the effect of intravenous thiamine on postoperative pain reduction in patients requiring maxillofacial surgery whom will be admitted in the Shiraz-chamran hospital. The following parameters such as: Individual patient information, medical history, the patient's hemodynamic values before- during and after surgery, duration of operation will be recorded. Severity of pain was evaluated by visual analogue scale (VAS) (100-mm) and the amount of opioid consumption after surgery, was recorded in maxillofacial ward. pain was measured after patient transferred to recovery room. (3) Setting and conduct: A total of 50 adult patients were selected for this randomized controlled trial. All subjects were divided into two groups to receive thiamine or placebo. While anesthesiologist, and student project manager would be unaware of an intravenous drugs. (

Medications taken by nurses are randomly coded ). An informed consent approved by the Ethics Committee of the Shiraz University of Medical Sciences. The anesthesiologist administered 200 mg of thiamine intravenously with 20 cc of 5% dextrose solution in study group and in the control group patients received placebo (IV) with 20 cc of 5% dextrose solution. Blood pressure and heart rate will be monitored and anesthetic drugs requirements during surgery, postoperative pain and necessity for postoperative opioids immediately after surgery, in the recovery room and until 12 hours after operation will be calculated and recorded. (4) Participants including major eligibility criteria: Inclusion criteria included 16-40year old patients whom were scheduled to undergo maxillofacial surgery! Subjects who had systemic disease and history of drug consumption, history of drug allergic reaction, hepatic and renal dysfunction and lack of signed informed consent will be excluded from the study sample . (5) Intervention: Patients in the study group will be received an intravenous "Thiamine hydrochloride" 200 mg dissolved in Dextrose Water 5% via pump infusion bolus for 15 min and the placebo group will receive " Dextrose Water 5% " as same bolus volume in a 15-minute intravenous infusion. (6) main outcome measures: During surgery heart rate, blood pressure and blood oxygen saturation is recorded at specified intervals and at the end of the operation analgesics usage were recorded. After surgery the patients were monitored for up to 4 hours in the recovery room and pain intensity (VAS 100-mm) is measured by a person who was uninformed about case and control groups every half hour. Pain relief drug consumption for patients will be recorded up to 12 hours in maxillofacial ward. post-operative complications will be asked in the questionnaire. Patient's satisfaction was evaluated as poor (0), moderate (1), good (2), and excellent (3).

#### General information

##### Acronym

## IRCT registration information

IRCT registration number: **IRCT201403061674N11**

Registration date: **2014-07-03, 1393/04/12**

Registration timing: **prospective**

Last update:

Update count: **0**

## Registration date

2014-07-03, 1393/04/12

## Registrant information

### Name

Hamid Reza Eftekharian

### Name of organization / entity

Shiraz University of Medical Sciences

### Country

Iran (Islamic Republic of)

### Phone

+98 71 3636 4001

### Email address

eftekhahr@sums.ac.ir

## Recruitment status

**Recruitment complete**

## Funding source

Supported by Vice-Chancellery of Research and Technology, Shiraz University Of Medical Sciences

## Expected recruitment start date

2014-08-23, 1393/06/01

## Expected recruitment end date

2015-07-21, 1394/04/30

## Actual recruitment start date

empty

## Actual recruitment end date

empty

## Trial completion date

empty

## Scientific title

Role of thiamine in the management of post-operative analgesia in maxillofacial surgery.

## Public title

Thiamine ( Intravenous Vitamin B1) induced post-operative analgesia in maxillofacial surgery.

## Purpose

Prevention

## Inclusion/Exclusion criteria

Inclusion criteria included: The patients with systemic disease and no history of drug use, and willingness to cooperate in this study are enrolled! Exclusion criteria included known allergy to the study drug, hepatic and renal dysfunction and lack of signed informed consent will be excluded.

## Age

From **16 years** old to **40 years** old

## Gender

Both

## Phase

2

## Groups that have been masked

*No information*

## Sample size

Target sample size: **50**

## 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

Ethics Committee of Shiraz University of Medical Sciences

##### Street address

Shiraz University of Medical Sciences, Vice-Chancellery of Research and Technology, Zand Avenue, Shiraz

##### City

Shiraz

##### Postal code

71345-1978

#### Approval date

2010-09-23, 1389/07/01

#### Ethics committee reference number

CT-P-93-7277

## Health conditions studied

### 1

#### Description of health condition studied

Maxillofacial surgery

#### ICD-10 code

00

#### ICD-10 code description

00

## Primary outcomes

### 1

#### Description

Intensity of postoperative pain

#### Timepoint

Immediately after surgery to 12 hours after surgery

#### Method of measurement

VAS

## 2

### **Description**

Use of narcotic after surgery

### **Timepoint**

To 12 hours after surgery

### **Method of measurement**

Nursing reports

## **Secondary outcomes**

### 1

### **Description**

post-operative complications of Thiamine

### **Timepoint**

Immediately after surgery to 12 hours after surgery

### **Method of measurement**

Nursing reports

### 2

### **Description**

Patient's level of satisfaction

### **Timepoint**

Immediately after surgery to 12 hours after surgery

### **Method of measurement**

poor (0), moderate (1), good (2), and excellent (3)

## **Intervention groups**

### 1

### **Description**

Patients in the interventional group (first group) will be received an intravenous "Thiamine hydrochloride" 200 mg dissolved in Dextrose Water 5% via pump infusion bolus for 15 min.

### **Category**

Treatment - Drugs

### 2

### **Description**

Patients in the the second group will receive " placebo" as same bolus volume of " Dextrose Water 5% " in a 15-minute intravenous infusion.

### **Category**

Treatment - Drugs

## **Recruitment centers**

### 1

### **Recruitment center**

#### **Name of recruitment center**

Shiraz Chamran Hospital

#### **Full name of responsible person**

Hamid Reza Eftekharian

#### **Street address**

Shiraz University Of Medical Sciences, Shiraz Chamran Hospital, Chamran Blvd., Shiraz

### **City**

Shiraz

## **Sponsors / Funding sources**

### 1

### **Sponsor**

#### **Name of organization / entity**

Vice-Chancellery of Research and Technology, Shiraz University Of Medical Sciences

#### **Full name of responsible person**

Sayed Basir Hashemi

#### **Street address**

Shiraz University of Medical Sciences, Vice-Chancellery of Research and Technology, Zand Avenue, Shiraz

#### **City**

Shiraz

#### **Grant name**

#### **Grant code / Reference number**

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

Yes

#### **Title of funding source**

Vice-Chancellery of Research and Technology, 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 Chamran Hospital

#### **Full name of responsible person**

Hamid Reza Eftekharian

#### **Position**

Assistant Professor

#### **Other areas of specialty/work**

#### **Street address**

Shiraz University Of Medical Sciences, Shiraz Chamran Hospital, Chamran Blvd., Shiraz

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**Person responsible for updating data****Contact****Name of organization / entity**

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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*