

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

11 Jul 2026

### **A comparison between the Effectiveness of two different Oral pastes, containing Triamcinolone and Licorice, on Minor Aphthous lesions: A triple blind randomized clinical trial**

#### **Protocol summary**

##### **Study aim**

Determining the Effectiveness of two types of oral pastes containing Licorice extract and Triamcinolone on oral Aphthous ulcers Using Herbal medicines (Licorice extraction) with fewer side effects to heal Aphthous lesions. Study the Effect of medicine on the size, pain and duration of lesion healing.

##### **Design**

A clinical trial has three groups receiving standard treatment, intervention and placebo or control group, triple blind, randomized, third phase on 60 patients, Using Balance block randomization

##### **Settings and conduct**

Patients who during the first 24 hours of the lesion appearing referred to Qazvin Dental School were examined by two oral medicine specialists and after confirming Aphthous lesions, Being informed and receiving consent, they entered one of the three groups completely randomly. The medicine containers will be the same and Coded and Given to the patient randomly by the secretary of the department, The examiner and the patient will be unaware of the contents of the medicine.

##### **Participants/Inclusion and exclusion criteria**

Inclusion criteria: No systemic disease (Hypertension, Diabetes, Hyperlipidemic, Bechet's disease, Rheumatoid disease) Not using medicines, cigarettes, alcohol and addiction. No-entry criteria: Allergic to herbal medicines. Oral ulcers that are not aphthous lesions. More than 48hrs passed from the incidence of the lesion. Use of medicines, cigarettes, alcohol and addiction. Systemic diseases

##### **Intervention groups**

Frist group (Standard treatment group): Using Oral paste containing Triamcinolone. The second group (Intervention group): Oral paste containing Licorice extract The third group (Control group): Placebo oral

paste without active substance

##### **Main outcome variables**

Size, Pain and Healing duration of the Lesion

#### **General information**

##### **Reason for update**

##### **Acronym**

##### **IRCT registration information**

IRCT registration number: **IRCT20210519051341N1**

Registration date: **2021-09-23, 1400/07/01**

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

Last update: **2021-09-23, 1400/07/01**

Update count: **0**

##### **Registration date**

2021-09-23, 1400/07/01

##### **Registrant information**

##### **Name**

masoume tork cheshme soltani

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

##### **Country**

Iran (Islamic Republic of)

##### **Phone**

+98 21 6678 5794

##### **Email address**

masoume.sln@gmail.com

##### **Recruitment status**

**Recruitment complete**

##### **Funding source**

##### **Expected recruitment start date**

2021-08-30, 1400/06/08

##### **Expected recruitment end date**

2021-09-30, 1400/07/08

##### **Actual recruitment start date**

empty

**Actual recruitment end date**

empty

**Trial completion date**

empty

**Scientific title**

A comparison between the Effectiveness of two different Oral pastes, containing Triamcinolone and Licorice, on Minor Aphthous lesions: A triple blind randomized clinical trial

**Public title**

A comparison between the Effectiveness of two different oral pastes, Containing Triamcinolone and Licorice, on minor Aphthous lesions: A triple blind randomized clinical trial

**Purpose**

Treatment

**Inclusion/Exclusion criteria****Inclusion criteria:**

Patients with Minor Aphthous ulcers who have been visited within the first 24 hours of incidence of the lesion. Patients do not have any systemic diseases ( Hypertension, Diabetes, Hyperlipidemia, Behcet disease, Rheumatism disease) Don't use medicine, cigarettes, alcohol and addiction

**Exclusion criteria:**

An oral ulcer that is not an Aphthous lesion More than 48hrs passed from incidence of the lesion Allergic to herbal medicines Use of medicine, cigarettes, alcohol and addiction Systemic diseases

**Age**

From **18 years** old to **65 years** old

**Gender**

Both

**Phase**

3

**Groups that have been masked**

- Participant
- Care provider
- Investigator
- Outcome assessor
- Data analyser
- Data and Safety Monitoring Board

**Sample size**

Target sample size: **60**

**Randomization (investigator's opinion)**

Randomized

**Randomization description**

Patients are divided into three treatment groups A, B, C using the Balance block randomization method, the size of each block is 6 and the total number of blocks is 10. Balanced randomization allocation method for participants, in a randomized clinical trial study of the effect of Oral paste, containing Triamcinolone (group A), placebo (group B) and Licorice(group C) in reducing the size, pain and healing duration of the lesion.

Randomization unit: individual Randomization tool: based on statistical software

**Blinding (investigator's opinion)**

Triple blinded

**Blinding description**

These patients are divided into three groups of triamcinolone, Licorice and placebo (10 blocks of 6) based on random allocation. Patients in the first group are given triamcinolone gel under the brand name Triadent 10 grams (from Raha Daroo Company) three times a day, the second group is given an oral gel containing Licorice and the third group is given a placebo gel. The containers in which the drugs are placed are very similar to each other and the contents are unknown. This study is triple blind and the patient, researcher and statistician are unaware of the type of drug which is given

**Placebo**

Used

**Assignment**

Parallel

**Other design features****Secondary Ids**

empty

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

Ethics committee of Qazvin University of Medical Sciences

**Street address**

Qazvin University of Medical Sciences, Bahonar Blv, Qazvin

**City**

Qazvin

**Province**

Qazvin

**Postal code**

3419915315

**Approval date**

2021-08-30, 1400/06/08

**Ethics committee reference number**

IR.QUMS.REC.1400.246

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

Recurrent oral Aphthae

**ICD-10 code**

K12.0

**ICD-10 code description**

Recurrent oral Aphthae

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

Amount of pain

### **Timepoint**

Before taking oral paste and the third, fifth and seventh days of treatment

### **Method of measurement**

On the visit days, we ask the patient to mark the amount of pain on the Visual Analogue Scale

## **2**

### **Description**

Healing duration

### **Timepoint**

Before taking oral paste and the third, fifth and seventh days of treatment

### **Method of measurement**

Days spent until complete healing of the lesion

## **3**

### **Description**

Size of the lesion

### **Timepoint**

Before taking oral paste and the third, fifth and seventh days of treatment

### **Method of measurement**

The largest diameter of the lesion is measured with the dental probe in millimetres

## **Secondary outcomes**

empty

## **Intervention groups**

### **1**

### **Description**

Intervention group: Patients in the intervention group receive a drug containing the Licorice active ingredient. They are trained to apply the gel on the Aphthous lesion 3 times a day after brushing and to avoid eating and drinking for half an hour after that. The drug consists of the Licorice active ingredient with 5% concentration and Carboxymethyl Cellulose (CMC).

### **Category**

Treatment - Drugs

### **2**

### **Description**

Standard treatment group: Triamcinolone (Trident 0.1%) is used as standard treatment. They should apply the oral paste on the Aphthous lesion 3 times a day after brushing and avoid eating and drinking for half an hour after that.

### **Category**

Treatment - Drugs

### **3**

### **Description**

Control group: This group receives a placebo (containing no active substance). The placebo only consists of

Carboxymethyl Cellulose (CMC). They should apply the placebo gel on the Aphthous lesion 3 times a day after brushing and avoid eating and drinking for half an hour after that.

### **Category**

Placebo

## **Recruitment centers**

### **1**

### **Recruitment center**

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

Department of Oral and maxillofacial medicine, school of dentistry, Qazvin University of Medical Sci

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

Mahdie Zarabadipour

#### **Street address**

Department of Oral and maxillofacial medicine, school of dentistry, Qazvin University of Medical Sciences, Bahonar Blv, Qazvin

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## **Sponsors / Funding sources**

### **1**

### **Sponsor**

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

Qazvin University of Medical Sciences

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

Doctor Seyed Mehdi Mirhashemi

#### **Street address**

Dept of research Qazvin University of Medical Sciences, Shahid Beheshti Ave, Qazvin, IRAN

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**Grant name**  
**Grant code / Reference number**  
**Is the source of funding the same sponsor organization/entity?**  
Yes  
**Title of funding source**  
Qazvin University of Medical Sciences  
**Proportion provided by this source**  
100  
**Public or private sector**  
Public  
**Domestic or foreign origin**  
Domestic  
**Category of foreign source of funding**  
*empty*  
**Country of origin**  
**Type of organization providing the funding**  
Academic

## Person responsible for general inquiries

### Contact

**Name of organization / entity**  
Qazvin University of Medical Sciences  
**Full name of responsible person**  
Mahdie Zarabadipour  
**Position**  
Associate professor, oral and maxillofacial medicine specialist  
**Latest degree**  
Specialist  
**Other areas of specialty/work**  
Dentistry  
**Street address**  
Department of Oral and maxillofacial medicine, school of dentistry, Qazvin University of Medical Sciences, Bahonar Blv, Qazvin  
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## Person responsible for scientific inquiries

### Contact

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## Person responsible for updating data

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## Sharing plan

### Deidentified Individual Participant Data Set (IPD)

Undecided - It is not yet known if there will be a plan to make this available

### Study Protocol

Undecided - It is not yet known if there will be a plan to make this available

### Statistical Analysis Plan

Undecided - It is not yet known if there will be a plan to make this available

**Informed Consent Form**

Undecided - It is not yet known if there will be a plan to make this available

**Clinical Study Report**

Undecided - It is not yet known if there will be a plan to make this available

**Analytic Code**

Undecided - It is not yet known if there will be a plan to make this available

**Data Dictionary**

Undecided - It is not yet known if there will be a plan to make this available