

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

10 Jun 2026

### Clinical trials the effect of aqueous extract of salvia officinalis for Preventing Stomatitis in Children With leukemia Undergoing Chemotherapy

#### Protocol summary

##### Summary

Objective: This single-blind clinical trial of the experimental and control groups with the aim of determine the effect of aqueous extract of sage on Preventing Stomatitis in children with Leukemia that admitted to the Hospital Besat Hematology Chemotherapy Unit in 1396. In this study, 54 patients with Cancer receiving chemotherapy were enrolled and randomly method available in any of the treatment groups (Rinse with water extract of sage) and control (Normal saline Mouthwash) are placed. In the control group patients only received routine treatment for patients with Stomatitis and in the experimental group used the routine treatment for mouth rinse in addition to 3 drops of extract in 2 ml water, diluted, enter the patient's mouth after brushing their teeth and mouth are rotated in and out for 1 minute and until 1 hour did not eat anything. To measure the effects of saline and aqueous extracts of sage on the severity of stomatitis use of the World Health Organization standard tools.

#### General information

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT2017011732025N1**

Registration date: **2017-02-10, 1395/11/22**

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

Last update:

Update count: **0**

##### Registration date

2017-02-10, 1395/11/22

##### Registrant information

##### Name

Arash Khalili

##### Name of organization / entity

Hamadan University Of Medical Sciences

##### Country

Iran (Islamic Republic of)

##### Phone

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##### Email address

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##### Recruitment status

**Recruitment complete**

##### Funding source

Hamadan University of Medical Sciences

##### Expected recruitment start date

2017-01-20, 1395/11/01

##### Expected recruitment end date

2017-05-22, 1396/03/01

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

##### Scientific title

Clinical trials the effect of aqueous extract of salvia officinalis for Preventing Stomatitis in Children With leukemia Undergoing Chemotherapy

##### Public title

The effect of aqueous extract of salvia officinalis for Preventing Stomatitis in Children With leukemia Undergoing Chemotherapy In Besat hospital.

##### Purpose

Prevention

##### Inclusion/Exclusion criteria

Inclusion criteria: The age of 6 to 14 years; Children's be admitted in Hematology Hospital Besat; children is leukemia; child or caregiver and are now teaching the

Persian language; baby have at least once chemotherapy and for the prevention of stomatitis to be hospitalized by doctor; Lack of sensitivity to drug sage Exclusion criteria: lack of desire to continue to participate in the study; out of the study for any reason during the study, such as death, relocation and ....

**Age**

From **5 years** old to **15 years** old

**Gender**

Both

**Phase**

N/A

**Groups that have been masked**

*No information*

**Sample size**

Target sample size: **70**

**Randomization (investigator's opinion)**

Randomized

**Randomization description****Blinding (investigator's opinion)**

Double blinded

**Blinding description****Placebo**

Not used

**Assignment**

Parallel

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

empty

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

Hamadan University Of Medical Scinces

**Street address**

Hamadan University Of Medical Scinces, Shahid Fahmideh Boulevard, Hamadan, Iran.

**City**

Hamadan

**Postal code****Approval date**

2017-01-14, 1395/10/25

**Ethics committee reference number**

IR.UMSHA.REC.1395.450

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

ALL

**ICD-10 code**

C91.0

**ICD-10 code description**

Acute lymphoblastic leukaemia [ALL]

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

Stomatitis

**Timepoint**

1week

**Method of measurement**

Oral Mucositis Assessment Scale

**Secondary outcomes****1****Description**

Oral Wound

**Timepoint**

1 Week

**Method of measurement**

Oral Mucositis Assessment Scale

**Intervention groups****1****Description**

This randomized, double-blind study, with a test group and a control group hematology hospital in Besat enforced. 60 leukemia patients undergoing chemotherapy simple random way in each of the treatment groups (rinse with water extract of sage and cocktails solution of 30 patients) and control (rinse with a solution of cocktails, 30) are placed. in the experimental group used the routine treatment for mouth rinse in addition to 3 drops of extract in 2 ml water, diluted, enter the patient's mouth after brushing their teeth and mouth are rotated in and out for 1 minute and until 1hour did not eat anything.

**Category**

Prevention

**2****Description**

In the control group patients only recived routine treatment for patients with Stomatitis

**Category**

Treatment - Drugs

**Recruitment centers****1****Recruitment center****Name of recruitment center**

Besat Hamadan Hospital

**Full name of responsible person**

Fariba Ebrahimi

**Street address**

Hamadan, Besat Hamadan Hospital

**City**  
Hamadan

## Sponsors / Funding sources

1

### Sponsor

**Name of organization / entity**  
Hamadan University Of Medical Sciences  
**Full name of responsible person**  
Fariba Ebrahimi Horyat  
**Street address**  
Hamadan University Of Medical Sciences  
**City**  
Hamadan

### Grant name

**Grant code / Reference number**

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

Yes

### Title of funding source

Hamadan 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

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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*