

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

09 Jun 2026

### Effect of propolis solution on prevention and treatment of oral mucositis in patients undergoing head and neck radiotherapy.

#### Protocol summary

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##### Summary

The goal of this randomized clinical trial is evaluation the effect of propolis solution on prevention and treatment of oral mucositis in patients undergoing the first episode of head and neck radiotherapy . This study is a randomized , double-blind and placebo- control. 60 patient who are referred to Imam Hossein Hospital for radiation therapy were randomly placed in two groups of 30 people (intervention and control group). The first group of patients will receive 5 drops of propolis solutionTDS (16 mg TDS ) for one month.The second group will received placebo. All patients will visit and evaluate weekly by radiotherapist and oral mucositis will record according to WHO, OMAS and NCI indices.

##### Recruitment status

**Recruitment complete**

##### Funding source

Pharmacy School of Shahid Beheshti University of Medical Sciences

##### Expected recruitment start date

2016-06-04, 1395/03/15

##### Expected recruitment end date

2017-06-05, 1396/03/15

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

#### General information

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT2016010225726N2**

Registration date: **2016-05-18, 1395/02/29**

Registration timing: **prospective**

Last update:

Update count: **0**

##### Registration date

2016-05-18, 1395/02/29

##### Registrant information

###### Name

Farzaneh Dastan

###### Name of organization / entity

Shahid Beheshti University of Medical Sciences

###### Country

Iran (Islamic Republic of)

###### Phone

+98 912 270 5933

###### Email address

##### Scientific title

Effect of propolis solution on prevention and treatment of oral mucositis in patients undergoing head and neck radiotherapy.

##### Public title

Effect of propolis solution on prevention and treatment of oral mucositis in patients undergoing head and neck radiotherapy.

##### Purpose

Prevention

##### Inclusion/Exclusion criteria

Inclusion criteria: patients undergoing the first episode of head and neck radiotherapy without any oral mucositis.Exclusion criteria: Chronic liver failure (Stage II & III); Chronic kidney disease (Stage IV & V); oral active infection; active collagen vascular disease; presence of any type of allergy and sensitivity to honey and propolis.

##### 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)

Double blinded

## Blinding description

## Placebo

Used

## Assignment

Parallel

## Other design features

## Secondary Ids

empty

## Ethics committees

### 1

#### Ethics committee

##### Name of ethics committee

Shahid Beheshti University of Medical Sciences

##### Street address

Tehran, Valiasr Street intersection of Niayesh Highway

##### City

Tehran

##### Postal code

#### Approval date

2016-03-01, 1394/12/11

#### Ethics committee reference number

IR.SBMU.PHNM.1394.309

## Health conditions studied

### 1

#### Description of health condition studied

oral mucositis

#### ICD-10 code

K12.3

#### ICD-10 code description

موکوزیت دهانی ناشی از رادیوتراپی

## Primary outcomes

### 1

#### Description

Mucositis severity

#### Timepoint

weekly

#### Method of measurement

Based on the criteria of NCI, OMAS & WHO

## Secondary outcomes

### 1

#### Description

Pain

#### Timepoint

Once a week for a month

#### Method of measurement

Based on the pain scale grade classification(0-10)

### 2

#### Description

Dysphagia

#### Timepoint

Once a week for a month

#### Method of measurement

Clinical evaluation and Questionnaire

## Intervention groups

### 1

#### Description

Intervention: administration of propolis , oral solution ,5 dropsTDS(16mg TDS), duration of intervention is one month

#### Category

Prevention

### 2

#### Description

Control group receives placebo

#### Category

Placebo

## Recruitment centers

### 1

#### Recruitment center

##### Name of recruitment center

Imam Hossein Hospital

##### Full name of responsible person

##### Street address

Tehran, Shahid madani street

##### City

Tehran

## Sponsors / Funding sources

### 1

#### Sponsor

##### Name of organization / entity

Shahid Beheshti University of Medical Sciences

##### Full name of responsible person

Farzaneh Dastan

##### Street address

Tehran-Valiasr Street intersection Niayesh Highway

**City**  
Tehran

**Grant name**

**Grant code / Reference number**

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

**Title of funding source**  
Shahid Beheshti 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**  
Shahid Beheshti University of Medical Sciences

**Full name of responsible person**  
Samaneh Dodge

**Position**  
PharmD. Student of Pharmacy

**Other areas of specialty/work**

**Street address**  
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## Person responsible for scientific inquiries

### Contact

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Farzaneh Dastan

**Position**

Assistant professor, Clinical pharmacy specialist

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## Person responsible for updating data

### Contact

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Shahid Beheshti University of Medical Sciences

**Full name of responsible person**  
Farzaneh Dastan

**Position**  
Assistant professor Clinical pharmacy specialist

**Other areas of specialty/work**

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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*