

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

30 May 2026

### Effect of Dexamethasone soaked nasalpack on postoperative pain of patients undergoing Dacryocystorhinostomy surgery

#### Protocol summary

##### Study aim

Determining the effect of topical dexamethasone on postoperative pain in patients under dacryocystorhinostomy

##### Design

Clinical trial with control group, with parallel groups, double-blind, randomized, phase 3 on 112 patients. Randaomaize.com was used for randomization.

##### Settings and conduct

In Khatam Al-Anbia Medical Center in Mashhad, after obtaining the patient's informed consent, they are randomly divided into two groups of intervention and control using computerized randomization software. For the intervention group, at the end of the operation, a dexamethasone-impregnated tampon is placed in the upper horn of the middle concha. In the control group, a tampon impregnated with normal saline is placed in the same place. Patients are asked in both groups in recovery and after entering the wards 3, 6, 12, 18 and 24 hours about their pain intensity with a verbal rating scale and also the amount of analgesic if it is used by persistent pain.

##### Participants/Inclusion and exclusion criteria

Inclusion criteria: Age between 18 and 75 years and able to speak Persian, Patient candidate for dacryocystorhinostomy surgery, Patients with class 1 and 2 anesthesia (ASA), no history of known and confirmed psychological disease, no history of current seizures, no history of alcohol or drug abuse based on history, no use of any analgesics in the 24 hours before surgery. Exclusion criteria: Any unusual complication during surgery (such as heavy bleeding during surgery, prolongation of surgery time (more than 75 minutes) and cardiopulmonary resuscitation during and after surgery

##### Intervention groups

For the intervention group, at the end of the operation, a dexamethasone-impregnated tampon is placed in the upper horn of the middle concha. In the control group, a tampon impregnated with normal saline is placed in the

same place.

##### Main outcome variables

Pain in the first 24 hours after surgery

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20210113050021N1**

Registration date: **2021-02-02, 1399/11/14**

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

Last update: **2021-02-02, 1399/11/14**

Update count: **0**

##### Registration date

2021-02-02, 1399/11/14

##### Registrant information

##### Name

Sara Jamali

##### Name of organization / entity

##### Country

Iran (Islamic Republic of)

##### Phone

+98 51 3728 9911

##### Email address

ocean\_jamali@yahoo.com

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2021-01-09, 1399/10/20

##### Expected recruitment end date

2022-01-10, 1400/10/20

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

**Trial completion date**  
empty

**Scientific title**  
Effect of Dexamethasone soaked nasalpack on postoperative pain of patients undergoing Dacryocystorhinostomy surgery

**Public title**  
Effect of Dexamethasone soaked nasalpack on postoperative pain of patients undergoing Dacryocystorhinostomy surgery

**Purpose**  
Treatment

**Inclusion/Exclusion criteria**  
**Inclusion criteria:**  
Age between 18 and 75 years old and able to speak Persian, patient candidate for dacryocystorhinostomy surgery, patients with class 1 and 2 anesthesia (ASA), no history of systemic hypertension based on history, no history of known and confirmed Psychological disease based on history, no history of current seizures based on history, no history of alcohol or drug abuse, no use of any painkillers in 24 hours before surgery  
**Exclusion criteria:**

**Age**  
From **18 years** old to **75 years** old

**Gender**  
Both

**Phase**  
3

**Groups that have been masked**

- Participant
- Outcome assessor

**Sample size**  
Target sample size: **112**

**Randomization (investigator's opinion)**  
Randomized

**Randomization description**  
Simple personal randomization will be done using random numbers table and www.randomization.com website. Randomization concealment will be done using closed envelopes. The envelopes will be prepared and printed by one of the members of the research team and random numbers with the help of Randomization.com and will be placed inside the envelope. The envelopes are closed and the contents cannot be seen from the outside. Then, the purpose of the study will be explained to the person who meets the inclusion criteria and if the person wishes and signs the informed consent form, takes an envelope and then opens it and enters the intervention or control group based on the contents of the envelope.

**Blinding (investigator's opinion)**  
Double blinded

**Blinding description**  
Patients and outcome evaluator are blinded in the trial about how patients are allocated.

**Placebo**  
Used

**Assignment**  
Parallel

**Other design features**

## Secondary Ids

empty

## Ethics committees

### 1

#### Ethics committee

##### Name of ethics committee

Ethics Committee of Mashhad University of Medical Sciences

##### Street address

Mashhad University of Medical Sciences, Azadi Ave., Mashhad

##### City

Mashhad

##### Province

Razavi Khorasan

##### Postal code

9177948564

#### Approval date

2020-11-09, 1399/08/19

#### Ethics committee reference number

IR.MUMS.MEDICAL.REC.1399.517

## Health conditions studied

### 1

#### Description of health condition studied

Lacrimal duct obstruction

#### ICD-10 code

H04.559

#### ICD-10 code description

Acquired stenosis of unspecified nasolacrimal duct

## Primary outcomes

### 1

#### Description

Pain in the first 24 hours after surgery

#### Timepoint

0, 3, 6, 12, 18 and 24 hours after surgery

#### Method of measurement

Pain intensity is assessed by the grading scale used in the NHS system of Wales University Hospital. This scale has 4 degrees: if the patient expresses complete pain relief, the pain intensity is zero and if the patient expresses severe pain, the patient's pain intensity is 3. At this scale, the patient's pain intensity is asked and the patient's description of the pain experienced is marked in one of four categories: "painless, mild, moderate, severe."

## Secondary outcomes

ocean\_jamali@yahoo.com

### 1

#### Description

Dosage in mg of rescue analgesia (morphine / pethidine / methadone / apotel / ketorolac / indomethacin)

#### Timepoint

0, 3, 6, 12, 18 and 24 hours after surgery

#### Method of measurement

questionnaire

### 2

#### Description

The interval between the first use of adjuvant painkillers and the end of the operation

#### Timepoint

In the first 24 hours after surgery

#### Method of measurement

Patient statement

## Intervention groups

### 1

#### Description

Intervention group: In the intervention group, at the end of the operation, a dexamethasone-impregnated tampon is inserted in the upper horn of the middle conch.

#### Category

Treatment - Drugs

### 2

#### Description

Control group: In the control group, at the end of the operation, a tampon impregnated with normal saline is inserted in the upper horn of the middle concha.

#### Category

Treatment - Drugs

## Recruitment centers

### 1

#### Recruitment center

##### Name of recruitment center

Khatam Al-Anbia Eye Hospital

##### Full name of responsible person

Sara Jamali

##### Street address

Khatam Al-Anbia Eye Hospital, Aboutaleb Intersection

##### City

Mashhad

##### Province

Razavi Khorasan

##### Postal code

9195965919

##### Phone

+98 51 3728 9911

##### Email

## Sponsors / Funding sources

### 1

#### Sponsor

##### Name of organization / entity

Mashhad University of Medical Sciences

##### Full name of responsible person

Khalil Abnous

##### Street address

Ghoreyshi building, Daneshgah Ave.

##### City

Mashhad

##### Province

Razavi Khorasan

##### Postal code

9195965919

##### Phone

+98 51 3845 2474

##### Email

ramresearch@mums.ac.ir

#### Grant name

#### Grant code / Reference number

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

Yes

#### Title of funding source

Mashhad 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

Mashhad University of Medical Sciences

##### Full name of responsible person

Sara Jamali

##### Position

Resident of Ophthalmology

##### Latest degree

Medical doctor

##### Other areas of specialty/work

Ophthalmology

##### Street address

Khatam Al-Anbia Eye Hospital, Aboutaleb Intersection

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**Person responsible for scientific inquiries**

**Contact**

**Name of organization / entity**

Mashhad University of Medical Sciences

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**Position**

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**Latest degree**

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**Person responsible for updating data**

**Contact**

**Name of organization / entity**

Mashhad University of Medical Sciences

**Full name of responsible person**

Sara Jamali

**Position**

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**Sharing plan**

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

Yes - There is a plan to make this available

**Study Protocol**

No - There is not a plan to make this available

**Statistical Analysis Plan**

No - There is not a plan to make this available

**Informed Consent Form**

No - There is not a plan to make this available

**Clinical Study Report**

No - There is not a plan to make this available

**Analytic Code**

No - There is not a plan to make this available

**Data Dictionary**

No - There is not a plan to make this available

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

Individual Participant Data Set (IPD): all collected deidentified IPD could be shared.

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

Starting 6 months after publication

**To whom data/document is available**

People working in academic institutions

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

No further criteria.

**From where data/document is obtainable**

: Khatam al-Anbia Eye Hospital, Aboutaleb Intersection, Mashhad, Iran. 00985137289911 - ocean\_jamali@yahoo.com

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

With a reasonable request, the data would be shared via email.

**Comments**