

Clinical Trial Protocol

Iranian Registry of Clinical Trials

10 Jun 2026

Investigating the Influence of Dry Eye Disease and Artificial Tear eye drop on the Biometry Repeatability and Accuracy in Patients with Cataract

Protocol summary

Study aim

Determining the Influence of Dry Eye Disease and Artificial Tear eye drop on the Biometry Repeatability and Accuracy in Patients with Cataract in the ophthalmology clinic of Farshchian (Sina) Hospital in 2024-2025

Design

Clinical trial in a group of 20 bilateral cataract surgery candidates (40 eyes) with dry eye disease referred to Hamedan Sina Hospital in 2024-25, with a control group, with parallel groups, not randomized and without blinding

Settings and conduct

In patients aged 50 to 70 years with bilateral cataracts, routine preoperative examinations will be performed. Schirmer test, TBUT test, and corneal staining will be used to determine the presence and severity of dry eye disease. Biometry will be performed twice using IOLMaster 500, 15 minutes apart, for one eye without artificial tears and ten minutes after applying artificial tear drop in the opposite eye, and this will be repeated. All patients will undergo Phaco cataract surgery under local anesthesia, and one month after surgery, patients' refraction will be measured and their spherical equivalent will be compared with the expected one.

Participants/Inclusion and exclusion criteria

Inclusion criteria: Patients aged 50 to 70 years with bilateral cataracts and mild to moderate dry eye disease without any pathology affecting biometry or prior ocular surgery and with consent to participate in the study
Exclusion criteria: Any complications during or after surgery or failure to attend follow-up visits

Intervention groups

Patients aged 50 to 70 years with bilateral cataracts and mild to moderate dry eye disease will be instilled with artificial tear drop in one eye before biometry, and the patient's refraction will be measured and the patients' SE will be compared with the expected SE in both eyes one month later.

Main outcome variables

Axial length; Anterior chamber depth; Corneal astigmatism; Mean keratometry; Spherical equivalent

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20250714066483N1**

Registration date: **2025-10-30, 1404/08/08**

Registration timing: **retrospective**

Last update: **2025-10-30, 1404/08/08**

Update count: **0**

Registration date

2025-10-30, 1404/08/08

Registrant information

Name

Elham Soltani

Name of organization / entity

Country

Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2024-12-21, 1403/10/01

Expected recruitment end date

2025-10-07, 1404/07/15

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date
empty

Scientific title
Investigating the Influence of Dry Eye Disease and Artificial Tear eye drop on the Biometry Repeatability and Accuracy in Patients with Cataract

Public title
Investigating the Influence of Dry Eye Disease and Artificial Tear eye drop on the Biometry Repeatability and Accuracy in Patients with Cataract

Purpose
Prevention

Inclusion/Exclusion criteria
Inclusion criteria:
Patients aged 50 to 70 years with bilateral cataract and mild to moderate Dry eye disease Informed consent to participate in the study Absence of pathologies influencing biometry such as keratoconus, corneal scar, pathologic myopia, nanophthalmus, etc Absence of prior ocular surgical history
Exclusion criteria:
Any intra- or postoperative complication occurrence Patient's failure to attend follow-up visit

Age
From **50 years** old to **70 years** old

Gender
Both

Phase
N/A

Groups that have been masked
No information

Sample size
Target sample size: **20**
More than 1 sample in each individual
Number of samples in each individual: **2**
both eyes in a patient with bilateral cataract and dry eye disease

Randomization (investigator's opinion)
Not randomized

Randomization description

Blinding (investigator's opinion)
Not blinded

Blinding description

Placebo
Not used

Assignment
Parallel

Other design features

Secondary Ids
empty

Ethics committees

1

Ethics committee
Name of ethics committee

Ethics committee of Hamedan university of medical sciences

Street address

Hamedan University of Medical Sciences, Shahid Fahmid Blvd., Pajouhesh Square

City

Hamedan

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Hamadan

Postal code

6517838736

Approval date

2025-02-16, 1403/11/28

Ethics committee reference number

IR.UMSHA.REC.1403.837

Health conditions studied

1

Description of health condition studied

cataract

ICD-10 code

H26

ICD-10 code description

Other cataract

2

Description of health condition studied

Dry eye disease

ICD-10 code

H04.12

ICD-10 code description

Dry eye syndrome

Primary outcomes

1

Description

spherical equivalent

Timepoint

At the beginning of the study and one month after surgery

Method of measurement

Topcon RM8000 autorefractometer

2

Description

Anterior chamber depth

Timepoint

At the beginning of the study and one month after surgery

Method of measurement

IOL Master 500

3

Description

Axial length

Timepoint

At the beginning of the study and one month after surgery

Method of measurement

IOL Master 500

4**Description**

mean keratometry

Timepoint

At the beginning of the study and one month after surgery

Method of measurement

Topcon RM8000 autorefractometer

5**Description**

corneal astigmatism

Timepoint

At the beginning of the study and one month after surgery

Method of measurement

Topcon RM8000 autorefractometer

6**Description**

Dry eye disease severity

Timepoint

before intervention

Method of measurement

Schirmer test, TBUT test, and corneal staining

Secondary outcomes

empty

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

Intervention group: primary biometry using IOLMaster-500 and then Instilling tearlose (Sinadaro) artificial tear drop 10 minutes before second biometry in one of a patient's eyes and one month after cataract surgery, measuring refraction and SE using auto refractometer

Category

Prevention

2**Description**

Control group: primary biometry using IOLMaster-500 and then not instilling Tearlose (Sinadaro) artificial tear drop 10 minutes before second biometry in one of a patient's eyes and one month after cataract surgery, measuring refraction and SE using auto refractometer

Category

Prevention

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

Sina (Farshchian) hospital

Full name of responsible person

Elham Soltani

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Sponsors / Funding sources**1****Sponsor****Name of organization / entity**

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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

Hamedan 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

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Person responsible for general inquiries**Contact****Name of organization / entity**

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Position

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Latest degree

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Hamadan

Province

Hamadan

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