Does silicone oil change ocular temperature?

Submission date 19/01/2017	Recruitment status No longer recruiting	 Prospectively registered Protocol
Registration date 30/01/2017	Overall study status Completed	 Statistical analysis plan Results
Last Edited 09/02/2017	Condition category Eye Diseases	 Individual participant data Record updated in last year

Plain English summary of protocol

Background and study aims

Eyes contain a jelly-like liquid called vitreous. When a patient is having surgery on their retina (the part of the eye where light signals are sent to the brain to create an image) this liquid has to be removed in order to gain access. This surgical procedure is known as a plars plana vitrectomy. During this procedure, vitreous needs to be replaced with a saline (salt water). In some cases, the retina can become detached from the eye and so silicone oil tamponade (gel) may be used to keep the retina in place. Silicone oil carries heat differently to water and these heat changes can lead to blurred vision. This study aims to compare eye temperature in participants who have an eye with silicone oil tamponade and an eye without it, in order to determine if silicone oil affects the temperature.

Who can participate?

Adults who are having a pars plana vitrectomy using silicone oil tamponade as part of their normal care.

What does the study involve?

The temperature of each participant's cornea (the out layer of the eye) is measured using a handheld non-contact thermometer in both eyes. This involves shining a special light into the eye which measures the temperature. Three temperature readings from each eye are taken. Participants also have their overall body temperature (taken from their forehead) measured using the same method.

What are the possible benefits and risks of participating? There are no direct benefits or risks to those taking part in the study.

Where is the study run from? Istanbul Education and Research Hospital (Turkey)

When is the study starting and how long is it expected to run for? November 2015 to May 2016

Who is funding the study? Investigator initiated and funded (Turkey) Who is the main contact? Dr. Erkan Unsal erkanunsal@gmail.com

Contact information

Type(s) Public

Contact name Dr Erkan Unsal

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers N/A

Study information

Scientific Title Temperature effect of silicone oil at the anterior segment

Study objectives The aim of this study is to measure the temperature difference between eyes with and without silicone oil tamponade.

Ethics approval required Old ethics approval format

Ethics approval(s)

Ethical Review Board of Istanbul Education and Research Hospital, 20/11/2015, ref: 2015-728

Study design

Observational single-centre cross-sectional study

Primary study design Observational

Secondary study design

Cross sectional study

Study setting(s)

Hospital

Study type(s)

Screening

Participant information sheet

Not available in web format, please use the contact details below to request a participant information sheet

Health condition(s) or problem(s) studied

Pars plana vitrectomy, intraocular silicone oil tamponade

Interventions

Participants have their central corneal eye temperature and their body temperature measured in a room with fixed temperature, humidity and illumination. To avoid inter-observer variations, all temperature measurements is done by a single examiner in the same examination room.

Central corneal temperatures are measured using a non-contact infrared thermometer from the corneal surface. The aiming beam of handheld infrared skin thermometer is centered slightly inferotemporally to the center of cornea. The working distance is the distance where the aiming beam formed a sharp concentric rings on the iris. Three measurements from each eye are taken and mean value for each eye is used for analysis.

Body temperature is measured using a non-contact infrared thermometer taken from the forehead. This device measures body temperature from 16 °C to 40 °C with a minimum accuracy of 0.2 °C. Measurements are obtained in less than one seconds in each patient.

Intervention Type

Other

Primary outcome measure

1. Corneal eye temperature both with and without silicone oil is measured using with noncontact infrared thermometer on the study visit

2. Body temperature is measured using a noncontact infrared thermometer on the study visit

Secondary outcome measures

No secondary outcome measures.

Overall study start date 20/11/2015

Completion date 01/09/2016

Eligibility

Key inclusion criteria 1. Require a pars plana vitrectomy and intraocular silicone oil tamponade 2. Aged between 36-88

Participant type(s) Patient

Age group Adult

Sex Both

Target number of participants 38 Male / 22 Female

Key exclusion criteria 1. Active ocular or periocular infection 2. Systemic infective illnesses

Date of first enrolment 01/12/2015

Date of final enrolment 30/05/2016

Locations

Countries of recruitment Türkiye

Study participating centre Istanbul Education and Research Hospital Department of Ophthalmology Kasap Ilyas Mah Org Abdurrahman Nafiz Gurman Cad Fatih Istanbul Türkiye 34098

Sponsor information

Organisation

Istanbul Research and Training Hospital

Sponsor details

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Sponsor type Hospital/treatment centre

ROR https://ror.org/00nwc4v84

Funder(s)

Funder type Not defined

Funder Name Investigator initiated and funded

Results and Publications

Publication and dissemination plan Not provided at time of registration.

Intention to publish date 01/09/2017

Individual participant data (IPD) sharing plan

The current data sharing plans for the current study are unknown and will be made available at a later date.

IPD sharing plan summary Data sharing statement to be made available at a later date