# Phase I trial: Ocular Technology Group International CV23-38

Submission date	Recruitment status	Prospectively registered
20/11/2024	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
26/11/2024	Deferred	Results
Last Edited	Condition category	Individual participant data
26/11/2024	Eye Diseases	<ul><li>Record updated in last year</li></ul>

## Plain English summary of protocol

The Health Research Authority (HRA) has approved deferral of publication of the full details of the trial. The full details will be added to the study record within 30 months after the trial has ended.

# **Contact information**

## Type(s)

Scientific, Principal investigator

#### Contact name

**Prof Michel Guillon** 

#### Contact details

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# Type(s)

Public

#### Contact name

Miss Deborah Moore

#### Contact details

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# Additional identifiers

#### Clinical Trials Information System (CTIS)

Nil known

## Integrated Research Application System (IRAS)

342156

#### ClinicalTrials.gov (NCT)

Nil known

#### Protocol serial number

ID23-37 CV23-38

# Study information

#### Scientific Title

Phase I trial: Ocular Technology Group International CV23-38 [The full scientific title will be published within 30 months after the end of the trial]

#### Study objectives

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# Ethics approval required

Ethics approval required

# Ethics approval(s)

approved 13/06/2024, South East Scotland Research Ethics Committee 02 (2nd Floor, Waverley Gate 2-4 Waterloo Place, Edinburgh, EH1 3EG, United Kingdom; +44 131 5369000; ruth. fraser4@nhslothian.scot.nhs.uk), ref: 24/SS/0049

# Study design

Non dispensing bilateral randomized study

# Primary study design

Interventional

# Study type(s)

Other

# Health condition(s) or problem(s) studied

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#### **Interventions**

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#### **Intervention Type**

Device

#### Phase

Phase I

## Drug/device/biological/vaccine name(s)

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#### Primary outcome(s)

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#### Key secondary outcome(s))

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## Completion date

29/10/2024

# **Eligibility**

# Key inclusion criteria

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# Participant type(s)

Healthy volunteer

# Healthy volunteers allowed

No

# Age group

Adult

# Lower age limit

40 years

#### Upper age limit

90 years

#### Sex

All

#### Total final enrolment

11

#### Key exclusion criteria

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#### Date of first enrolment

01/08/2024

#### Date of final enrolment

25/10/2024

# Locations

#### Countries of recruitment

**United Kingdom** 

England

# Study participating centre

Ocular Technology Group International

66 Buckingham Gate LONDON United Kingdom SW1E 6AU

# **Sponsor information**

#### Organisation

CooperVision International Limited

# Funder(s)

# Funder type

Industry

#### Funder Name

CooperVision International Ltd.

# **Results and Publications**

# Individual participant data (IPD) sharing plan

The datasets generated and/or analysed during the current study are not expected to be made available because of their high commercial sensitivity and the negligible benefit to the public of publication of results of non-therapeutic clinical trials.

# IPD sharing plan summary

Not expected to be made available

#### **Study outputs**

Output type Details Date created Date added Peer reviewed? Patient-facing?

Participant information sheet Participant information sheet 11/11/2025 11/11/2025 No Yes