

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

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

## 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

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### Type(s)

Public

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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?
<a href="#">Participant information sheet</a>	Participant information sheet	11/11/2025	11/11/2025	No	Yes