

Deferred registration - Quotient Sciences code: QSC303694

Submission date 04/06/2026	Recruitment status Not yet recruiting	<input checked="" type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 24/06/2026	Overall study status Deferred	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 24/06/2026	Condition category Other	<input type="checkbox"/> Individual participant data <input checked="" type="checkbox"/> Record updated in last year

Plain English summary of protocol

This is a phase I trial in healthy volunteers only and qualifies for an automatic deferral. The full details will be added to the trial record on or before the deferral expiry date.

Contact information

Type(s)

Principal investigator

Contact name

Dr David Everton

Contact details

Mere Way, Ruddington Fields
Ruddington
Nottingham
United Kingdom
NG11 6JS
+44 (0)1159749000
recruitment@weneedyou.co.uk

Type(s)

Scientific, Public

Contact name

None Clinical Operations

Contact details

Avenue de la Gare 39
Lausanne
Switzerland
1003

+41 (0)21 711 3970
info@oculis.com

Additional identifiers

Integrated Research Application System (IRAS)
1013199

CRO Study Code
QSC303694

Study information

Scientific Title

CTIMP - Quotient Sciences code: QSC303694 [The full scientific title will be added on or before the deferral expiry date]

Study objectives

This is a phase I trial in healthy volunteers only and qualifies for an automatic deferral. The full details will be added to the trial record on or before the deferral expiry date.

Ethics approval required

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Ethics approval(s)

Approved 08/06/2026, London – Harrow Research Ethics Committee (Health Research Authority 2 Redman Place Stratford, London, E20 1JQ, United Kingdom; +44 (0)2071048357; harrow.rec@hra.nhs.uk), ref: 26/LO/0330

Primary study design

Interventional

Allocation

N/A: single arm study

Masking

Open (masking not used)

Control

Uncontrolled

Assignment

Single

Purpose

Phase I study in healthy volunteers

Study type(s)

Health condition(s) or problem(s) studied

Healthy volunteers

Interventions

This is a phase I trial in healthy volunteers only and qualifies for an automatic deferral. The full details will be added to the trial record on or before the deferral expiry date.

Intervention Type

Drug

Phase

Phase I

Drug/device/biological/vaccine name(s)

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Primary outcome(s)

1. [Outcome name] measured using [Metric or method of measurement] at [Timepoint(s)]. This is a phase I trial in healthy volunteers only and qualifies for an automatic deferral. The full details will be added to the trial record on or before the deferral expiry date.

Key secondary outcome(s)

Completion date

12/08/2026

Eligibility

Key inclusion criteria

This is a phase I trial in healthy volunteers only and qualifies for an automatic deferral. The full details will be added to the trial record on or before the deferral expiry date.

Healthy volunteers allowed

Yes

Age group

Mixed

Lower age limit

18 Years

Upper age limit

65 Years

Sex

Male

Total final enrolment

0

Key exclusion criteria

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Date of first enrolment

06/07/2026

Date of final enrolment

12/08/2026

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Quotient Sciences Limited

Mere Way, Ruddington Fields

Ruddington

Nottingham

England

NG11 6JS

Sponsor information

Organisation

Neurocol Operations Sàrl

Funder(s)

Funder type

Funder Name

Neurocol Operations Sàrl

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not expected to be made available