Phase I trial, Quotient Code: QSC300720

Submission date	Recruitment status	[X] Prospectively registered
14/07/2023	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
19/07/2023	Deferred	Results
Last Edited	Condition category	Individual participant data
19/07/2023	Other	Record updated in last year

Plain English summary of protocol

The sponsor has confirmed that the trial meets the criteria for 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)

Principal investigator

Contact name

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Type(s)

Scientific

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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)

1007785

ClinicalTrials.gov (NCT)

Nil known

Protocol serial number

IRAS 1007785

Study information

Scientific Title

Phase I trial, Quotient Code: QSC300720 [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)

1. submitted 06/07/2023, London Surrey Borders REC (2nd Floor 2 Redman Place, Stratford, London, E20 1JQ, United Kingdom; +44 (0)207 104 8057; Surreyborders.rec@hra.nhs.uk), ref: 23 /LO/0521

2. submitted 06/07/2023, MHRA (10 South Colonnade, Canary Wharf, London, E14 4PU, United Kingdom; +44 (0)20 3080 6000; info@mhra.gov.uk), ref: CTA 58311/0001/001-0001

Study design

One-part single-centre non-randomized open-label 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

Drug

Phase

Phase I

Drug/device/biological/vaccine name(s)

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Primary outcome(s)

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Key secondary outcome(s))

The sponsor has confirmed that the trial meets the criteria for 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.

Completion date

Eligibility

Key inclusion criteria

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Participant type(s)

Healthy volunteer

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

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Date of first enrolment

27/09/2023

Date of final enrolment

16/02/2024

Locations

Countries of recruitment

United Kingdom

England

Study participating centre Quotient Sciences Limited

Mere Way Ruddington Fields Ruddington Nottingham United Kingdom NG11 6JS

Sponsor information

Organisation

Veradermics Inc

Funder(s)

Funder type

Industry

Funder Name

Veradermics Inc

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 No Yes