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Phase 1 Trial: Quotient Code QSC303303, Sponsor Code LMNL6511C1002

Submission date 22/04/2025	**Recruitment status** Recruiting	[X] Prospectively registered [_] Protocol [_] Statistical analysis plan [_] Results [_] Individual participant data [X] Record updated in last year
Registration date 02/05/2025	**Overall study status** Deferred	
Last Edited 16/05/2025	**Condition category** Other	

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. Full details will be added to the study record within 30 months after the trial has ended.

Contact information

Type(s)
Public, Scientific

Contact name
Dr Clinical Trial Information

Contact details
Liminal BioSciences Limited, 3rd Floor, 1 Ashley Road
Altrincham
United Kingdom
WA14 2DT
-
info@liminalbiosciences.com

Type(s)
Principal investigator

Contact name
Dr Nand Singh

Contact details
Quotient Sciences Limited, Mere Way, Ruddington Fields
Nottingham
United Kingdom
NG11 6JS

+44 (0)330 303 1000
recruitment@weneedyou.co.uk

Additional identifiers

Clinical Trials Information System (CTIS)
Nil known

Integrated Research Application System (IRAS)
1011852

ClinicalTrials.gov (NCT)
Nil known

Protocol serial number
Quotient Code QSC303303, Sponsor Code LMNL6511C1002

Study information

Scientific Title
Phase 1 Trial: Quotient Code QSC303303, Sponsor Code LMNL6511C1002

Study objectives
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Ethics approval required
Ethics approval required

Ethics approval(s)
approved 14/05/2025, London - Surrey Borders Research Ethics Committee (Equinox House, City Link, Nottingham, NG2 4LA, United Kingdom; +44 (0)20 7104 8057; surreyborders.rec@hra.nhs.uk), ref: 25/LO/0200

Study design
Two part single centre randomized double-blind study to assess biomarkers safety tolerability and pharmacokinetics

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. Full details will be added to the study record within 30 months after the trial has ended.

Completion date
04/05/2026

Eligibility

Key inclusion criteria
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Participant type(s)
Healthy volunteer

Healthy volunteers allowed
No

Age group
Adult

Lower age limit
18 years

Upper age limit
65 years

Sex
All

Key exclusion criteria
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Date of first enrolment
02/06/2025

Date of final enrolment
04/05/2026

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
Liminal BioSciences Limited

Funder(s)

Funder type
Industry

Funder Name
Liminal BioSciences Limited

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