ISRCTN90651766 https://doi.org/10.1186/ISRCTN90651766

Phase I trial: Quotient code QSC301813

Submission date 31/07/2024	**Recruitment status** No longer recruiting	[X] Prospectively registered [_] Protocol [_] Statistical analysis plan [_] Results [_] Individual participant data [_] Record updated in last year
Registration date 02/08/2024	**Overall study status** Deferred	
Last Edited 25/09/2024	**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. The full details will be added to the study record within 30 months after the trial has ended.

Contact information

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Additional identifiers

Integrated Research Application System (IRAS)
1009723

Protocol serial number
Quotient code: QSC301813

Study information

Scientific Title
Phase I trial: Quotient code QSC301813

Study objectives
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Ethics approval required
Ethics approval required

Ethics approval(s)
Approved 13/09/2024, London – Brent Research Ethics Committee (Health Research Authority, 2 Redman Place, Stratford, London, E20 1JQ, United Kingdom; +44 (0)207 104 8117; brent.rec@hra.nhs.uk), ref: 24/LO/0348

Primary study design
Interventional

Study design
Relative bioavailability study in 14 healthy volunteers

Study type(s)
Other, Safety

Health condition(s) or problem(s) studied
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Interventions
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.

Intervention Type
Drug

Phase
Phase I

Drug/device/biological/vaccine name(s)
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Primary 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.

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
29/01/2025

Eligibility

Key inclusion criteria
Healthy volunteer

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
16/09/2024

Date of final enrolment

29/01/2025

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
Quotient Sciences Limited
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Ruddington Fields
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Sponsor information

Organisation
Carmot Therapeutics (United States)

ROR
https://ror.org/05ye86j67

Funder(s)

Funder type
Industry

Funder Name
Carmot Therapeutics, Inc.

Results and Publications

Individual participant data (IPD) sharing plan
The datasets generated during 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