ISRCTN13656987 https://doi.org/10.1186/ISRCTN13656987

Deferred Registration: Quotient Sciences Code: QSC302324

Submission date 19/05/2026	**Recruitment status** Recruiting	[X] Prospectively registered [_] Protocol [_] Statistical analysis plan [_] Results [_] Individual participant data [X] Record updated in last year
Registration date 13/07/2026	**Overall study status** Deferred	
Last Edited 13/07/2026	**Condition category** Other	

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

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Type(s)
Public, Scientific

Contact name
Dr Clinical Trials Information

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Additional identifiers

Integrated Research Application System (IRAS)
1010387

Study information

Scientific Title
Deferred Registration: Quotient Sciences Code: QSC302324 [Full scientific title to 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
Ethics approval required

Ethics approval(s)
Approved 29/10/2024, Health and Social Care Research Ethics Committee A (HSC REC A) (Lissue Industrial Estate West, 5 Rathdown Walk, Lisburn, BT28 2RF, United Kingdom; +44 2895361400; reca@hscni.net), ref: 24/NI/0088

Primary study design
Interventional

Study design
Two-part, single centre, randomised, first-in-man study to assess safety, pharmacokinetics, pharmacodynamics and food effect in 84 healthy volunteers.

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)
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)
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

Completion date
20/11/2027

Eligibility

Key inclusion criteria
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Participant type(s)
Healthy volunteer

Healthy volunteers allowed
Yes

Age group
Adult

Sex
All

Total final enrolment
0

Key exclusion criteria
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Date of first enrolment
17/11/2026

Date of final enrolment
20/11/2027

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
Quotient Sciences Limited
Mere Way
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Sponsor information

Organisation
Pheno Therapeutics Limited

Funder(s)

Funder type
Industry

Funder Name
Pheno Therapeutics Limited

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not expected to be made available