ISRCTN13906399 https://doi.org/10.1186/ISRCTN13906399

Phase 1 trial HMR code: 25-006

Submission date 25/09/2025	**Recruitment status** No longer recruiting	[X] Prospectively registered [_] Protocol
Registration date 29/09/2025	**Overall study status** Deferred	[_] Statistical analysis plan [_] Results
Last Edited 23/09/2026	**Condition category** Other	[_] Individual participant data [X] 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.

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Additional identifiers

Integrated Research Application System (IRAS)
1012527

HMR code
25-006

Sponsor code
MED-TSN-102

Study information

Scientific Title
Phase 1 trial HMR code: 25-006
The full scientific title will be published within 30 months after the end of the trial

Study objectives
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.

Ethics approval required
Ethics approval required

Ethics approval(s)
Approved 01/09/2025, London – Brent Research Ethics Committee (2 Redman Place, Stratford, London, E20 1JQ, United Kingdom; +44 20 7104 8128; brent.rec@hra.nhs.uk), ref: 25/LO/0462

Primary study design
Interventional

Study design
Safety and pharmacokinetics trial in up to 36 healthy women

Study type(s)
Other

Health condition(s) or problem(s) studied
Healthy volunteers

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)
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.

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
31/03/2027

Eligibility

Key inclusion criteria
Healthy human female volunteers

Participant type(s)
Healthy volunteer

Healthy volunteers allowed
No

Age group

Mixed

Sex
Female

Total final enrolment
0

Key exclusion criteria
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.

Date of first enrolment
07/10/2025

Date of final enrolment
21/09/2026

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
Hammersmith Medicines Research (HMR)
Cumberland Avenue, Park Royal
London
England
NW10 7EW

Sponsor information

Organisation
Medherant Limited

Funder(s)

Funder type
Industry

Funder Name
Medherant Limited

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not expected to be made available