ISRCTN60181900 https://doi.org/10.1186/ISRCTN60181900

Phase 1 trial: Fortrea CRU code: 8526072

Submission date 04/08/2025	**Recruitment status** No longer recruiting	[ ] Prospectively registered [ ] Protocol [ ] Statistical analysis plan [ ] Results [ ] Individual participant data [ ] Record updated in last year
Registration date 04/08/2025	**Overall study status** Deferred	
Last Edited 04/08/2025	**Condition category** Other	

Plain English summary of protocol
The Health Research Authority (HRA) has approved 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)
1009878

Study information

Scientific Title
Phase 1 trial: Fortrea CRU code: 8526072

Acronym
Nil known

Study objectives
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Ethics approval required
Ethics approval required

Ethics approval(s)
Approved 30/10/2024, North East REC – York (2 Redman Place, London, E20 1JQ, United Kingdom; +44 207 104 8052; york.rec@hra.nhs.uk), ref: 24/NE/0157

Primary study design
Interventional

Study design
Interventional

Study type(s)
Other

Health condition(s) or problem(s) studied
The Health Research Authority (HRA) has approved 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.

Interventions
The Health Research Authority (HRA) has approved 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
Other

Primary outcome(s)

The Health Research Authority (HRA) has approved 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 Health Research Authority (HRA) has approved 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
02/03/2025

Eligibility

Key inclusion criteria
Healthy human volunteer

Participant type(s)
Healthy volunteer

Healthy volunteers allowed
No

Age group
Adult

Sex
Male

Key exclusion criteria
Does not meet inclusion criteria

Date of first enrolment
29/01/2025

Date of final enrolment
04/02/2025

Locations

Countries of recruitment
United Kingdom

England

Study participating centre

Fortrea Clinical Research Unit Limited
Draper's Yard Marshall Street
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United Kingdom
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Sponsor information

Organisation
CellCentric Limited

Funder(s)

Funder type
Industry

Funder Name
CellCentric 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 publication of results of non-therapeutic clinical trials.

IPD sharing plan summary
Not expected to be made available