

A Phase 1 trial: MEU Phase 1 Clinic: SUDO-286-102

Submission date 05/09/2025	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 08/09/2025	Overall study status Deferred	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 08/09/2025	Condition category Other	<input type="checkbox"/> Individual participant data <input checked="" type="checkbox"/> Record updated in last year

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

Type(s)

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Additional identifiers

Clinical Trials Information System (CTIS)

Nil known

Integrated Research Application System (IRAS)

1006866

ClinicalTrials.gov (NCT)

Nil known

Protocol serial number

Nil known

Study information

Scientific Title

A Phase 1 trial: MEU Phase 1 Clinic: SUDO-286-102

Study objectives

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Ethics approval required

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Ethics approval(s)

approved 20/06/2024, North West - Greater Manchester Central Research Ethics Committee (2 Redman Place, Stratford, London, E20 1JQ, United Kingdom; +44 2071048023; gmcentral.rec@hra.nhs.uk), ref: 24/NW/0062

Study design

Interventional single-centre blinded randomized study

Primary study design

Other

Study type(s)

Safety

Health condition(s) or problem(s) studied

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Interventions

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Intervention Type

Other

Primary outcome(s)

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Key secondary outcome(s)

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Completion date

17/04/2025

Eligibility

Key inclusion criteria

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Participant type(s)

Healthy volunteer, Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

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Date of first enrolment

20/06/2024

Date of final enrolment

18/12/2024

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Medicines Evaluation Unit Limited

The Langley Building

Southmoor Road

Wythenshawe

Manchester

United Kingdom

M23 9QZ

Sponsor information

Organisation

Sudo Biosciences

Funder(s)

Funder type

Industry

Funder Name

Sudo Biosciences

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