

Phase I trial; HMR code: 24-003

Submission date 05/08/2024	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 12/08/2024	Overall study status Deferred	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 06/01/2025	Condition category Other	<input type="checkbox"/> Individual participant data <input type="checkbox"/> 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.

Contact information

Type(s)

Principal investigator

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Additional identifiers

Clinical Trials Information System (CTIS)

Nil known

Integrated Research Application System (IRAS)

1010220

Protocol serial number

IRAS 1010220; HMR code: 24-003; Sponsor code: SLG-TED-101

Study information

Scientific Title

Phase I trial; HMR code: 24-003 [The full scientific title will be published within 30 months after the end of the trial]

Study objectives

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Ethics approval required

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Ethics approval(s)

approved 30/07/2024, London – Surrey Borders Research Ethics Committee (Equinox House, City Link, Nottingham, NG2 4LA, United Kingdom; +44 207 104 8057; surreyboundaries.rec@hra.nhs.uk), ref: 24/LO/0492

Study design

Pharmacokinetic and bioavailability study in up to 20 healthy volunteers

Primary study design

Interventional

Study type(s)

Other

Health condition(s) or problem(s) studied

Healthy volunteers

Interventions

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Intervention Type

Drug

Phase

Phase I

Drug/device/biological/vaccine name(s)

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Primary outcome(s)

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Key secondary outcome(s)

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Completion date

08/08/2025

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

14/08/2024

Date of final enrolment

08/05/2025

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Hammersmith Medicines Research Limited

Cumberland Avenue

London

United Kingdom

NW10 7EW

Sponsor information

Organisation

Sling Therapeutics, Inc.

Funder(s)

Funder type

Industry

Funder Name

Sling Therapeutics, Inc.

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