ISRCTN17405388 https://doi.org/10.1186/ISRCTN17405388

Phase I trial BDD code: BDD22320

Submission date 24/07/2025	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol
Registration date 24/07/2025	**Overall study status** Deferred	[_] Statistical analysis plan [_] Results
Last Edited 24/07/2025	**Condition category** Other	[_] Individual participant data [X] 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 12 months after the trial has ended.

Contact information

Type(s)
Public, Scientific, Principal investigator

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Additional identifiers

Clinical Trials Information System (CTIS)
Nil known

Integrated Research Application System (IRAS)
1009022

ClinicalTrials.gov (NCT)
Nil known

Protocol serial number
MOS118-101

Study information

Scientific Title
Phase I trial BDD code: BDD22320

Study objectives
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Ethics approval required
Ethics approval required

Ethics approval(s)
approved 04/03/2024, London - Brent Research Ethics Committee (2 Redman Place, Stratford, London, E20 1JQ, United Kingdom; +44 (0)2071048128; brent.rec@hra.nhs.uk), ref: 24/LO/0003

Study design
Pharmacoscintigraphic open-label crossover study

Primary study design
Interventional

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)
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Key secondary outcome(s))
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Completion date
18/08/2025

Eligibility

Key inclusion criteria
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Participant type(s)
Healthy volunteer

Healthy volunteers allowed
No

Age group
Adult

Lower age limit
18 years

Upper age limit
65 years

Sex
All

Key exclusion criteria
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Date of first enrolment
09/06/2025

Date of final enrolment
23/06/2025

Locations

Countries of recruitment
United Kingdom

Scotland

Study participating centre
BDD Pharma Ltd
Glasgow Royal Infirmary
84 Castle Street
Glasgow
United Kingdom
G4 0SF

Sponsor information

Organisation
Mosanna Therapeutics

Funder(s)

Funder type
Industry

Funder Name
Mosanna Therapeutics

Results and Publications

Individual participant data (IPD) sharing plan
The datasets generated and/or analysed during the study are not expected to be made available because of the high commercial sensitivity of this phase I study and the negligible benefit to the public of phase I information.

IPD sharing plan summary
Not expected to be made available