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Multimodal Augmented Reality for Operative Guidance in Interventional Neuroradiology

Submission date 14/08/2024	**Recruitment status** Recruiting	[_] Prospectively registered [_] Protocol
Registration date 03/09/2024	**Overall study status** Deferred	[_] Statistical analysis plan [_] Results
Last Edited 16/05/2025	**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 30 months after the trial has ended.

Contact information

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Principal investigator

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Additional identifiers

Clinical Trials Information System (CTIS)
Nil known

Integrated Research Application System (IRAS)
319189

ClinicalTrials.gov (NCT)
Nil known

Protocol serial number
CPMS 54208

Study information

Scientific Title
Multimodal Augmented Reality for Operative Guidance in Interventional Neuroradiology

Acronym
MAROG-INR

Study objectives
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Ethics approval required
Ethics approval required

Ethics approval(s)
approved 21/02/2023, London - Dulwich Research Ethics Committee (Skipton House, 80 London Road, London, SE1 6LH, United Kingdom; +44 (0)207 104 8290; dulwich.rec@hra.nhs.uk), ref: 23/LO/0030

Study design
Single-centre interventional trial in patients undergoing interventional neuroradiology procedures

Primary study design

Interventional

Study type(s)
Treatment

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))
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
01/07/2026

Eligibility

Key inclusion criteria
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Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Adult

Sex
All

Key exclusion criteria
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Date of first enrolment
01/09/2024

Date of final enrolment
01/04/2026

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
Charing Cross Hospital
Fulham Palace Road
London
United Kingdom
W6 8RF

Sponsor information

Organisation
Medical iSight (UK) Limited

Funder(s)

Funder type
Other

Funder Name
Self-funded

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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes