

Multimodal Augmented Reality for Operative Guidance in Interventional Neuroradiology

Submission date 14/08/2024	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 03/09/2024	Overall study status Deferred	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 16/05/2025	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)

Principal investigator

Contact name

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Type(s)

Public, Scientific

Contact name

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Additional identifiers

Clinical Trials Information System (CTIS)

Nil known

Integrated Research Application System (IRAS)

319189

Protocol serial number

CPMS 54208

Study information

Scientific Title

Multimodal Augmented Reality for Operative Guidance in Interventional Neuroradiology

Acronym

MAROG-INR

Study objectives

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.

Ethics approval required

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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

Intentional

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)

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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