

# Multimodal Augmented Reality for Operative Guidance in Interventional Neuroradiology

<b>Submission date</b> 14/08/2024	<b>Recruitment status</b> Recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 03/09/2024	<b>Overall study status</b> Deferred	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
<b>Last Edited</b> 16/05/2025	<b>Condition category</b> 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

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Public, Scientific

### Contact name

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**Additional identifiers****EudraCT/CTIS number**

Nil known

**IRAS number**

319189

**ClinicalTrials.gov number**

Nil known

**Secondary identifying numbers**

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**

Interventional

## **Secondary study design**

### **Study setting(s)**

Hospital

### **Study type(s)**

Treatment

### **Participant information sheet**

Not available in web format

### **Health condition(s) or problem(s) studied**

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### **Interventions**

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### **Intervention Type**

Other

### **Primary outcome measure**

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### **Secondary outcome measures**

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### **Overall study start date**

29/11/2022

### **Completion date**

01/07/2026

## **Eligibility**

### **Key inclusion criteria**

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### **Participant type(s)**

Patient

**Age group**

Adult

**Sex**

Both

**Target number of participants**

50

**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**

England

United Kingdom

**Study participating centre**

**Charing Cross Hospital**

Fulham Palace Road

London

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## **Sponsor information**

**Organisation**

Medical iSight (UK) Limited

**Sponsor details**

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**Sponsor type**  
Industry

**Website**  
<https://www.medicalisight.com/>

## **Funder(s)**

**Funder type**  
Other

**Funder Name**  
Self-funded

## **Results and Publications**

### **Publication and dissemination plan**

Full trial details will be published up to 30 months after the end of the trial. Results will be posted on or after the date of publication of full trial details.

**Intention to publish date**  
21/02/2026

### **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