

Multimodal Augmented Reality for Operative Guidance in Oncoplastic Breast Surgery

Submission date 07/11/2024	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 11/11/2024	Overall study status Deferred	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 11/11/2024	Condition category Other	<input type="checkbox"/> Individual participant data <input 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

Mr Daniel Leff

ORCID ID

<https://orcid.org/0000-0002-5310-1046>

Contact details

Department of Breast Surgery

First Floor

Charing Cross Hospital

Fulham Palace Road

London

United Kingdom

W6 8RF

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d.leff@imperial.ac.uk

Type(s)

Public, Scientific

Contact name

Dr Philip Pratt

Contact details

Medical iSight (UK) Ltd
9th Floor
107 Cheapside
London
United Kingdom
EC2V 6DN

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pjp@medicalisight.com

Additional identifiers**Clinical Trials Information System (CTIS)**

Nil known

Integrated Research Application System (IRAS)

306799

ClinicalTrials.gov (NCT)

Nil known

Protocol serial number

IRAS 306799

Study information**Scientific Title**

Multimodal Augmented Reality for Operative Guidance in Oncoplastic Breast Surgery

Acronym

MAROG - OBS

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 10/02/2022, London - Queen Square Research Ethics Committee (RA NRES Centre Manchester, Barlow House, 3rd floor, 4 Minshull Street, Manchester, M1 3DZ, United Kingdom; +44 (0)207 104 8225; queensquare.rec@hra.nhs.uk), ref: 21/PR/1795

Study design

Multi-centre interventional non-control trial in patients undergoing oncoplastic breast procedures

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

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Interventions

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.

Intervention Type

Other

Primary outcome(s)

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Key secondary outcome(s)

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Completion date

31/12/2025

Eligibility**Key inclusion criteria**

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Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

Female

Key exclusion criteria

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Date of first enrolment

31/05/2022

Date of final enrolment

31/12/2025

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre**Charing Cross Hospital**

Department of Breast Surgery
1st Floor, Charing Cross Hospital
Fulham Palace Road
London
United Kingdom
W6 8RF

Study participating centre**St George's University Hospital**

Department of Plastic and Reconstructive Surgery
Blackshaw Road
Tooting
London
United Kingdom
SW17 0QT

Sponsor information**Organisation**

Imperial College London

ROR

<https://ror.org/041kmwe10>

Funder(s)

Funder type

Other

Funder Name

Investigator initiated and funded

Results and Publications

Individual participant data (IPD) sharing plan

Not expected to be made available.

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