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Multimodal Augmented Reality for Operative Guidance in Oncoplastic Breast Surgery

Submission date 07/11/2024	**Recruitment status** No longer recruiting	[ ] Prospectively registered [ ] Protocol [ ] Statistical analysis plan [ ] Results [ ] Individual participant data [ ] Record updated in last year
Registration date 11/11/2024	**Overall study status** Deferred	
Last Edited 11/11/2024	**Condition category** Other	

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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Additional identifiers

Clinical Trials Information System (CTIS)
Nil known

Integrated Research Application System (IRAS)
306799

Protocol serial number
IRAS 306799

Study information

Scientific Title
Multimodal Augmented Reality for Operative Guidance in Oncoplastic Breast Surgery

Acronym
MAROG - OBS

Study objectives
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Ethics approval required
Ethics approval required

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))
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
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
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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