

Testing to see how virtual reality mindfulness and wellbeing session before and after cardiac surgery will help to reduce anxiety and stress

Submission date 03/01/2024	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
Registration date 05/01/2024	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 06/10/2025	Condition category Surgery	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Background and study aims

Non-drug methods like hypnosis and Virtual Reality (VR) are used alongside other treatments to help patients with anxiety and different types of pain. But when it comes to using VR to help people who are having heart surgery, there haven't been many studies. One reason for this might be that using VR for this purpose is relatively new, VR technology is changing quickly, and it can be expensive to get started with VR.

This pilot study will assess immersive head-mounted VR simulations in adult patients undergoing first-time elective cardiac surgery to understand the impact on patient anxiety and stress. The primary objective is to assess anxiety and stress levels between two groups.

Who can participate?

Patients aged 18 years or older, undergoing cardiac surgery at Wythenshawe Hospital.

What does the study involve?

Patients will be randomly assigned to two groups: one receiving standard care (control group) and the other receiving VR intervention. The control group will go through the regular admission process, including informational resources and music sessions using a wireless headset. The VR group will receive the same, but with additional VR sessions before surgery and on the third day after surgery. We will exclude patients requiring ventilation on the third day and replace them with new participants in both groups. Both groups will complete questionnaires before and after surgery to assess anxiety and stress levels. Physiological assessments, VR experience, and absorption will be recorded for the VR group. Additionally, we will explore the relationship between salivary cortisol levels (a potential stress biomarker) and self-reported anxiety levels using questionnaires. Four sets of samples will be collected before and after the placebo/VR interventions to examine the correlation.

What are the possible benefits and risks of participating?

Benefits: The potential benefit if you are allocated to try the Virtual Reality immersive intervention is reduced stress, anxiety and improved wellbeing and mindfulness after using the

device.

Risks: Some people may experience nausea, dizziness, disorientation, balance problems, headache, eye strain or fatigue when using VR headsets, although this is rare. If you experience any of these symptoms, or feel uncomfortable in any way, you are free to remove the headset at any time and you do not need to carry on with the study.

In addition, there is a risk of cross-contamination of infection from one patient to another patient when using VR headsets and headphones. To avoid any possible cross-contamination, we will be cleaning the headphones and headsets with alcoholic disinfectant frequently.

Where is the study run from?
University of Salford (UK)

When is the study starting and how long is it expected to run for?
January 2024 to June 2024

Who is funding the study?
University of Salford (UK)

Who is the main contact?
Prof Bhuvanewari Krishnamoorthy, b.bibleraaj@salford.ac.uk

Contact information

Type(s)

Public, Scientific, Principal investigator

Contact name

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Additional identifiers

Clinical Trials Information System (CTIS)

Nil known

Integrated Research Application System (IRAS)

335136

Protocol serial number

IRAS 335136

Study information

Scientific Title

A Randomised pilot Study to evaluate the use of Virtual Reality Mindfulness and Wellbeing session during pre- and post-surgical admission to reduce anxiety and stress undergoing first time elective cardiac surgery – RSVR study

Acronym

RSVR

Study objectives

The use of Immersive head mounted Virtual Reality headset intervention can reduce pre-and postoperative anxiety and stress in adult patients undergoing elective cardiac surgery

Ethics approval required

Ethics approval required

Ethics approval(s)

approved 02/01/2024, North West - Greater Manchester East Research Ethics Committee (3rd Floor, Barlow House, 4 Minshull Street, Manchester, M1 3DZ, United Kingdom; +44 207 104 8290; gmeast.rec@hra.nhs.uk), ref: 23/NW/0345

Study design

Single-centre two-arm randomized controlled pilot study

Primary study design

Interventional

Study type(s)

Quality of life, Efficacy

Health condition(s) or problem(s) studied

Pre-and postoperative anxiety and stress in adult patients undergoing elective cardiac surgery

Interventions

Cardiac surgery patients (n = 30) undergoing first time elective/urgent-in-patient cardiac surgery at Wythenshawe Hospital site will be randomly assigned equally to two arms (control with placebo or VR) using a computer-generated randomisation list. The control group will receive standard care admission process which includes a surgical information booklet, online Microsoft Teams experiential talk about the surgery pathway by a heart charity and two sessions of 15 minutes music session with WorWoder Wireless blue tooth headset as a placebo pre- and post-surgery as same timings as intervention group. The VR intervention group will receive the same as the control group with the addition of two VR sessions: session one 15 minutes VR session will be one hour before surgery and sessions two on day 3 evening (around 6pm) postoperatively. Patients who are ventilated on day 3 will be removed from the study and additional patients will be recruited on both groups. Questionnaires will be handed to the patients pre- and post-surgery in both groups to rate their levels of anxiety and stress level

(STAI), physiological assessments, VR experience and VR-absorption (VAS) in intervention group will be recorded.

In addition, we will explore if there is association between salivary cortisol levels as a potential biomarker for stress and self-completed questionnaires anxiety level (STAI). We will obtain four samples of before and after placebo/VR intervention pre- and post-surgery to see the correlation.

Intervention Type

Device

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

VR experience from Rescape Innovations (<https://www.rescape.health/>)

Primary outcome(s)

Patients' anxiety and stress levels measured using STAI before and after each session on pre- and post-surgery

Key secondary outcome(s)

1. Cortisol swabs before & after each session on pre- and post-surgery
2. Physiological parameters pre- and post-surgery in each session:
 - 2.1. Heart rate measured using Holter monitor which is standard in all cardiac wards at the NHS hospital
 - 2.2. Blood pressure measured using Holter monitor which is standard in all cardiac wards at the NHS hospital
 - 2.3. Respiratory rate measured using direct observation by the researcher

Completion date

30/06/2024

Eligibility

Key inclusion criteria

1. Adults undergoing first-time elective cardiac surgery (coronary artery bypass surgery and aortic/mitral valve surgeries)
2. Age more than 18 years
3. Give informed consent for their participation to this study
4. Urgent inpatients who are well enough to read and understand the study and patient information leaflet

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Upper age limit

100 years

Sex

All

Total final enrolment

36

Key exclusion criteria

1. Psychiatric antecedents, claustrophobia or acrophobia, diagnosis or self-reported anxiety disorder, hearing or visual impairment, Infectious cases (example: conjunctivitis), redo surgery, critical preoperative state, impaired cognitive function, current diagnosis of epilepsy, dementia, or other neurologic disease, receipt of anxiolytic drugs or sedatives within the preceding 24 hours, or injury to the face, or non-infectious, infectious skin conditions that would prevent comfortable use of the VR hardware.
2. Patients who are having complex cardiac surgeries.
3. Patients who are admitted for an emergency cardiac surgery.
4. Patients who do not want to listen to music on their bedside Bluetooth enabled headphones.
5. Patients who do not want to wear VR headsets.

Date of first enrolment

05/02/2024

Date of final enrolment

30/06/2024

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

University Hospital of South Manchester NHS Foundation Trust

Wythenshawe Hospital

Southmoor Road

Wythenshawe

Manchester

United Kingdom

M23 9LT

Sponsor information

Organisation

University of Salford

ROR

<https://ror.org/01tmqtf75>

Funder(s)

Funder type

University/education

Funder Name

University of Salford Manchester

Alternative Name(s)

University of Salford, University of Salford, Manchester, USM

Funding Body Type

Private sector organisation

Funding Body Subtype

Universities (academic only)

Location

United Kingdom

Results and Publications

Individual participant data (IPD) sharing plan

All data will be anonymised. Patients who have consented to get a copy of the results will be posted to their home address.

The datasets generated during and/or analysed during the current study are available from the corresponding author on reasonable request and will be included in the subsequent results publication.

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IPD sharing plan summary

Available on request, Published as a supplement to the results publication

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article		04/10/2025	06/10/2025	Yes	No

version 0.2

Participant information sheet		12/12/2023	05/01/2024	No	Yes
Protocol file	version 0.1	03/10/2023	05/01/2024	No	No