

# Impact of a remote lifestyle coaching program on the length of hospital stay and complications in patients undergoing knee and hip arthroplasty surgery

<b>Submission date</b> 31/10/2022	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
<b>Registration date</b> 14/11/2022	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 15/03/2023	<b>Condition category</b> Musculoskeletal Diseases	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

## Plain English summary of protocol

### Background and study aims

Sapien Health is a mobile app-based behavioural intervention for patients undergoing elective surgery. The programme combines personalised educational programmes with remote health coaching to support patients to optimise their modifiable risk factors for surgery, and support their recovery during the postoperative phase.

This study aims to assess the impact of this app on individual and population health through the modification of unhealthy lifestyle behaviours for patients undergoing elective surgery for knee and hip arthroplasty. It is also hoped that by facilitating the delivery of remote perioperative care, Sapien Health can play a supportive role in the NHS response to the COVID-19 pandemic.

The Enhanced Recovery After Surgery model (ERAS) has been instrumental in emphasising the need for an end-to-end approach, and its implementation has seen improvements in both clinical and financial outcomes. Nevertheless, avoidable costs related to surgery remain high, with day-of-surgery cancellations estimated to cost the NHS £400m per year, and postoperative wound complications amounting to £980m per year.

Encouraging patients to be more active in their own care before and after surgery is increasingly recognised as an opportunity to improve outcomes and reduce some of these avoidable costs. This is especially true for patients undergoing major surgery, where the impact of modifiable lifestyle factors on surgical outcomes is particularly well-established. Increasingly, the time before and after surgery is seen as a “teachable moment” - a window of opportunity to address unhealthy behaviours, with consequent individual and population health benefits.

Traditional approaches to lifestyle modification using patient information leaflets and isolated educational interventions during outpatient clinic appointments are now understood to be largely ineffective. Reasons commonly cited for this include low health literacy and psychological reactance as a consequence of a failure to involve patients in the decision-making

process. For lifestyle interventions to be effective, a shared decision-making approach with active collaboration between patient and clinician is key.

Who can participate?

Adults aged 18 years old and over listed for elective knee and hip arthroplasty

What does the study involve?

The study will assess if using the app increases a patient's knowledge, skills and confidence in managing their own well-being. These will be measured using a questionnaire called PAM (Patient Activation Measure). Evidence shows that when people are supported to become more active, they benefit from better health outcomes, improved care experiences and fewer unplanned admissions. This study will also see if using the app helps to reduce patients' length of hospital stay following surgery, postoperative complications, re-admission to hospital following discharge and surgery cancellation rate. We will measure these things by randomly assigning half the patients who have chosen to participate in the study to use the app and comparing it against the PAM score and surgical outcome metrics of half the patients who will not use the app.

What are the possible benefits and risks of participating?

No benefits and risks were provided at registration.

Where is the study run from?

Sherwood Forest Hospitals NHS Foundation Trust, Kingsmill Hospital (UK)

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

December 2021 to October 2023

Who is funding the study?

National Institute for Health and Care Research (NIHR) (UK)

Who is the main contact?

Dr Rebecca Barker (UK)

rebecca.barker6@nhs.net

## Contact information

### Type(s)

Principal investigator

### Contact name

Dr Rebecca Barker

### Contact details

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## **Additional identifiers**

**Integrated Research Application System (IRAS)**  
311052

**Central Portfolio Management System (CPMS)**  
52124

## **Study information**

### **Scientific Title**

Impact of a remote lifestyle change program on length of stay and complications in patients undergoing elective knee and hip arthroplasty

### **Acronym**

SAPIEN SBRI 19

### **Study objectives**

As a solution to patients having a remote lifestyle, Sapien, a mobile app-based behavioural intervention for patients undergoing elective surgery, combines personalized digital guidance with 1-to-1 remote health coaching to help optimize patients preoperatively and support their recovery during the postoperative phase.

The app aims to modify risk by supporting patients to:

1. Increase physical activity levels
2. Stop smoking
3. Reduce alcohol intake
4. Improve diet
5. Improve sleep duration and quality
6. Enhance preparedness for their perioperative journey

The study will assess if using the app increases PAM (Patient Activation Measure) score, this is a questionnaire which measures a patient's knowledge, skills and confidence in managing their own wellbeing. Evidence shows that when people are supported to become more active, they benefit from better health outcomes, improved care experiences and fewer unplanned admissions. This study will also see if using the app helps to reduce patients' length of hospital stay following surgery, postoperative complications, re-admission to hospital following discharge and surgery cancellation rate. We will measure these things by randomly assigning half the patient population who have chosen to participate in the study use of the app and comparing it against the PAM score and surgical outcome metrics of half the population who will not use the app.

### **Ethics approval required**

Old ethics approval format

### **Ethics approval(s)**

Approved 17/03/2022, East Midlands - Nottingham 2 Research Ethics Committee (C/o: Carolyn Halliwell, Approvals Specialist, Equinox House, City Link, Nottingham, HRA1 Meeting Room, NG2 4LA, UK; +44 (0)207 1048098; nottingham2.rec@hra.nhs.uk), ref: 22/EM/0063

## **Study design**

Interventional randomized controlled multicentre study

## **Primary study design**

Interventional

## **Study type(s)**

Treatment

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

Behavioural intervention for patients undergoing primary lower limb arthroplasty

## **Interventions**

This clinical trial will randomise 200 patients to either the intervention arm (use of the app) or control (usual care.) A randomisation code will be computer generated, in permuted blocks of 10 patients and stratified according to age. The 100 patients in the interventional arm will use the app within their peri-operative management for at least 6 weeks pre-operative. The data collected from these patients will be compared to data from the 100 patients (comparison arm), for these patients only the primary outcome and secondary outcomes 1, 2, 3 and 4 will be collected by the clinical team.

The data collected by the clinical team from both arms will be anonymized before being provided to the Sapien team for analysis. Data collected by the Sapien application will be made available to the clinical team where applicable. The effects of the Sapien Health application on the primary outcome and secondary outcomes 1, 2, 3 and 4 will be evaluated by means of comparison between the 'interventional arm' and the 'comparison arm'.

## **Study Duration**

For the 'interventional arm' the Sapien Health application will be available for a maximum of 10 weeks prior to surgery and for 1 month after. The exact amount of time patients utilise the Sapien Health application prior to surgery will vary with a minimum of 6 weeks pre-surgery required to make effective use of the app. Study duration for the 'control arm' will be a maximum of 6 weeks prior to surgery and for up to 90 days post-operative to allow collection of outcomes from patients' medical notes.

## **Intervention Type**

Behavioural

## **Primary outcome(s)**

Intervention success measured using the Patient Activation Measure (PAM) scoring at programme entry and pre-operatively (before surgery) and up to 30 days post-surgery

## **Key secondary outcome(s)**

1. Postoperative complications within 30-days of surgery (pulmonary and cardiac) measured using medical records electronic and paper 30 days post-surgery
2. Readmission to the hospital within 30 days after surgery measured using an electronic database called CareFlow that sends notifications regarding research patient admissions at any

time from randomisation

3. Length of stay measured using medical records and the hospital database that records admission and discharge data

4. The proportion of cancellations of surgeries measured using data collected from randomised patients and recorded on the trial database. Collected by the trial team and analysed by Sapien Health.

5. Health behaviour change measured using questionnaires provided by Sapien and recorded on REDCap. Sapien health to then analyse.

### **Completion date**

01/10/2023

## **Eligibility**

### **Key inclusion criteria**

1. Adults aged 18 years old and over listed for elective knee and hip arthroplasty

2. Date of the surgery is (or is expected to be) a minimum of 6 weeks from the date of onboarding for the control and intervention arms

3. Access to a smartphone with Apple IOS or Android operating systems and willingness to install the Sapien Health application

4. Sufficient confidence in written and spoken English to provide informed consent and utilise the Sapien Health application

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

Patient

### **Healthy volunteers allowed**

No

### **Age group**

Adult

### **Lower age limit**

18 years

### **Sex**

All

### **Key exclusion criteria**

1. Individuals undergoing emergency surgery

2. Pregnant or breastfeeding individuals.

### **Date of first enrolment**

29/06/2022

### **Date of final enrolment**

31/05/2023

## **Locations**

**Countries of recruitment**

United Kingdom

England

**Study participating centre**

**Sherwood Forest Hospitals NHS Foundation Trust**

Kings Mill Hospital

Mansfield Road

Sutton-in-ashfield

United Kingdom

NG17 4JL

**Study participating centre**

**Royal Berkshire NHS Foundation Trust**

Royal Berkshire Hospital

London Road

Reading

United Kingdom

RG1 5AN

## Sponsor information

**Organisation**

Sherwood Forest Hospitals NHS Foundation Trust

**ROR**

<https://ror.org/04ce87537>

## Funder(s)

**Funder type**

Government

**Funder Name**

National Institute for Health and Care Research

**Alternative Name(s)**

National Institute for Health Research, NIHR Research, NIHRresearch, NIHR - National Institute for Health Research, NIHR (The National Institute for Health and Care Research), NIHR

## Funding Body Type

Government organisation

## Funding Body Subtype

National government

## Location

United Kingdom

# Results and Publications

## Individual participant data (IPD) sharing plan

1. Findings will be made available to Sherwood Forest Hospitals and Royal Berkshire Hospital for dissemination internally and any relevant hospital communications. Data will also be presented in lay language on the Sapien Health website.
2. The findings will be written up as a case study and Sapien Health will seek to publish in relevant academic journals and present at conferences.
3. The datasets, including the type of surgery, surgery date, postoperative complications, readmission to hospital, email address, questionnaire results and trial numbers, will be available 30 days post last patient recruited from Luke Eastwood, luke@sapienhealth.io (Sapien Health). Consent from participants was required and obtained. Data were anonymized and will be identified by the trial ID given on REDCap.

## IPD sharing plan summary

Available on request

## Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">HRA research summary</a>			28/06/2023	No	No