

Feasibility study of hybrid or cemented implants for total hip replacement

Submission date 30/10/2020	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
Registration date 19/02/2021	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 10/12/2025	Condition category Surgery	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Current plain English summary as of 02/03/2022:

Background and study aims

The hip joint is made up of a ball and socket joint, with the ball being at the head of the thigh bone (femur) and the cup-shaped socket in the pelvic bone. A total hip replacement involves devices being surgically implanted to replace damaged bone within the hip joint and this study is interested in two of the most common types. One type is where both the stem in the thigh bone and the socket in the pelvic bone are cemented in place and the other is where the stem is cemented and the socket is uncemented (called a hybrid implant). Both types of implant are regularly used in similar numbers but little research had been done to establish if one is any better than the other. The aim of this small study is to see if patients who are undergoing hip replacement for osteoarthritis are willing to participate in a study to provide information on how well they consider their new hip to be functioning, and how it affects their day-to-day activities, so that the researchers can decide if a larger study will be achievable to see which implant is better.

Who can participate?

Patients aged 18 and above who are undergoing total hip replacement and for whom either of the study implants are suitable to be used (fully cemented or hybrid)

What does the study involve?

Participants will have the type of implant used selected at random and the researchers will collect information about the operation from clinical notes. At recruitment and 6 weeks and between 3-6 months after their operation participants will be asked to answer questionnaires about the effect their hip replacement has on their day-to-day activity and whether they have had to use NHS services. In addition to this, a mix of people made up of those who took part and those who declined to take part, as well as surgeons and healthcare professionals involved in recruitment and/or data collection, will be interviewed by researchers to learn how recruitment and data collection processes may be developed to improve these aspects in a future trial.

An analysis of cost to the NHS for the treatment of the arthritis and the hip replacement procedure itself will also be carried out. This will be done by comparing what health services (e.g. visits to GP) were required for the six months before and after the surgery.

What are the possible benefits and risks of participating?

The possible risks and benefits of taking part are no different to usual clinical care as the hip implants and study procedures are in line with routine care. Participants will be asked to complete a short set of questionnaires on three occasions (at recruitment and 6 weeks and between 3-6 months post-operation) and some participants, surgeons and healthcare professionals may also take part in interviews with researchers.

Where is the study run from?

Liverpool Clinical Trials Centre (UK)

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

May 2020 to April 2022

Who is funding the study?

National Institute for Health Research (NIHR) (UK)

Who is the main contact?

Tony Coffey

hiphop@liverpool.ac.uk

Previous plain English summary:

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Who is the main contact?
Tony Coffey
hiphop@liverpool.ac.uk

Contact information

Type(s)
Scientific

Contact name
Mr Tony Coffey

Contact details
Trial Coordinator
Liverpool Clinical Trials Centre
University of Liverpool
Block C, Waterhouse Building
1-5 Brownlow Street
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United Kingdom
L69 3GL
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hiphop@liverpool.ac.uk

Additional identifiers

Clinical Trials Information System (CTIS)
Nil known

Integrated Research Application System (IRAS)
271885

ClinicalTrials.gov (NCT)
Nil known

Protocol serial number
CPMS 46257, IRAS 271885

Study information

Scientific Title

Feasibility study for a comparative trial of hybrid or cemented implants for total hip replacement. Hip arthroplasty with Hybrid Or cemented implants: Patient-reported outcomes (HipHOP)

Acronym

HipHOP

Study objectives

The aim of this study is to establish the feasibility of performing a randomized controlled trial comparing patient-reported functional outcomes and cost-effectiveness of total hip arthroplasty using either a fully cemented or a hybrid total hip implant.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approved 26/08/2020, HSC REC B (Office for Research Ethics Committees Northern Ireland (ORECNI), Customer Care & Performance Directorate, Lissue Industrial Estate West, 5 Rathdown Walk, Moira Road, Lisburn, BT28 2RF, UK; +44 (0)28 95361400 ; recb@hscni.net), ref: 20/NI/0096

Study design

Randomized; Both; Design type: Treatment, Device, Surgery, Qualitative

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Total hip replacement for osteoarthritis

Interventions

Current intervention as of 02/03/2022:

The design of this study incorporates both quantitative and qualitative data collection and analyses, referred to as Workstream 1 (WS1) and Workstream 2 (WS2), respectively. Workstream 1 is the randomized controlled trial comparing the two hip implants by Patient-Reported Outcome Measures (PROMS); a cost-utility analysis is also included in the PROMS. Workstream 2 is interviews with patients, surgeons and healthcare professionals, designed to obtain their thoughts on the randomized controlled trial. The study has been discussed with a PPI group.

All patients that are approached, regardless of their decision to participate in Workstream 1, will also be invited to participate in Workstream 2 if they meet the eligibility criteria, on which further information is shown below.

Those patients that participate in workstream one will be allocated at random to receive either a fully cemented hip joint (where both the stem and socket are cemented into place) or a hybrid

hip joint (where just the stem is cemented and the body forms new bone to hold the socket). They will also be required to complete a portfolio of questionnaires (validated and bespoke) on three occasions. There are six questionnaires in total, though not all of them need to be completed on every occasion.

The first occasion must be prior to them being randomized to the type of hip joint they will receive. If they complete the questionnaires on the day of their surgery, it must also be prior to them receiving any anaesthetic.

The second occasion will be at their routine 6-week postoperative follow-up and the third occasion will be via post at between 3-6 months post operation. If the routine 6-week post-operative consultation is virtual (e.g. due to COVID-19), the questionnaires will be posted to the patient at that timepoint too.

Patients will not need to make any extra visits to hospital than they would for standard care and each will be on study for between 3-6 months post-operatively.

Workstream two will aim to recruit up to 30 patients, 20-30 surgeons and about 6 healthcare professionals to participate in interviews.

Patient interviews will be conducted to gain an understanding of why they chose to participate in workstream one, or not, along with their experiences of being in the study or their perceptions of what it would entail and, in the case of decliners, what the barriers were that prevented them from doing taking part. These interviews will be conducted over the phone. For participants who also take part in workstream 1, these interviews will ideally take place between 2-4 weeks after the patient has undergone their surgery. For decliners, the interview will be arranged at a time convenient to the participant, which could be before or after surgery.

Surgeons that practice at hospitals participating in Workstream 1 (both surgeons who are, and who are not, allowing their patients to be randomised to Workstream 1), and also at other sites that could be considered as recruiting centres for a larger trial will be approached to participate in the workstream two interviews. Initial approach will be made by the Chief Investigator. He will provide a participant information sheet to explain the study and introduce the qualitative researcher, based at the University of Manchester. The purpose of these interviews will be to understand their perceptions of the trial and equipoise, understand factors relating to willingness/unwillingness to take part in a trial, and to identify any other issues that could affect trial findings being implemented into practice. These interviews will be done over the phone.

The third cohort of participants for Workstream 2 are healthcare professionals who approached patients to participate in workstream 1 of the study and/or were involved in data collection. The purpose of these interviews is to learn about their experiences in recruiting patients and collecting data, and to learn about any changes to the study that may be required to aid recruitment to the larger study. These interviews will be done over the phone.

Previous interventions as of 27/07/2021:

The design of this study incorporates both quantitative and qualitative data collection and analyses, referred to as Workstream 1 (WS1) and Workstream 2 (WS2), respectively. Workstream 1 is the randomized controlled trial comparing the two hip implants by Patient-Reported Outcome Measures (PROMS); a cost-utility analysis is also included in the PROMS. Workstream 2 is interviews with patients, surgeons and healthcare professionals, designed to obtain their thoughts on the randomized controlled trial. The study has been discussed with a PPI group.

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Patients will not need to make any extra visits to hospital than they would for standard care and will each be on study for 6 months post-operatively (-1 month/+2 months).

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Intervention Type

Procedure/Surgery

Primary outcome(s)

Current primary outcome measure as of 27/07/2021:

Recruitment rate measured by the:

1. Total number of patient participants randomised per month
2. Ratio of successful recruitment to eligible patients approached

Timepoint(s): 7 months

Previous primary outcome measure:

Recruitment rate measured by the:

1. Total number of patient participants randomised per month
2. Ratio of successful recruitment to eligible patients approached

Timepoint(s): 12 months

Key secondary outcome(s)

Current secondary outcome measures as of 27/07/2021:

1. Adherence to the protocol will be measured by:

- 1.1. The number of minor (e.g. visit time-point violation) or major (e.g. violation of inclusion criteria) protocol deviations collected on a patient participant and site level between randomisation and the defined end of study
- 1.2. The percentage of patients in each arm that are accidentally unblinded measured between randomisation and the defined end of study

2. Trial withdrawal rate at 6 weeks and 3-6 months will be measured using the total number of patients randomised in the study. In addition, withdrawal rates will be presented by trial withdrawal reasons and withdrawal stage for both arms.

3. Patient participant population characteristics will be measured by collecting the following information:

- 3.1. Age – at randomisation
- 3.2. BMI – at randomisation
- 3.3. Proportion of missing data at the point of final analysis
- 3.4. Loss to follow up at the point of final analysis

4. Clinical and patient-reported data will be measured by collecting the following information:

- 4.1. Patient participant-reported outcomes including change in OHS, FJS, WPAI-SHP, SAPS score and quality of life measured using the EQ-5D- 5LTM at 6 weeks and 3-6 months
- 4.2. Revision rate, defined as the number of patient participants needing revision arthroplasty within 3-6 months of surgery
- 4.3. Infection rate, measured as the rate of re-operation for infection within 3-6 months of surgery
- 4.4. Length of stay, measured in hours between patient admission and initial discharge
- 4.5. Operation time, measured in minutes as the differences between the start and end of the operation
- 4.6. Incidence of treatment cross over measured as the surgically reported incidence of a patient receiving an intervention other than the one they were allocated through the randomisation system.
- 4.7. Incidence of intra-operative surgical complications, the proportion of patient participants

experiencing greater than grade III severity will be measured and reported according to Clavien Dindo classification system during patient follow-up

4.8. Incidence of postoperative surgical complications, the proportion of patient participants experiencing greater than grade III severity will be measured and reported according to Clavien Dindo classification system during patient follow-up

5. Completeness of the EQ-5D-5L (Trademark) and Healthcare Resource Use Survey will be measured by using the following methods:

5.1. Descriptive statistics will be used to assess the frequency of missing data for the EQ-5D-5L (Trademark) and Healthcare Resource Use Survey at the point of final analysis.

5.2. Any patterns in missing data will be identified through the use of descriptive statistics at the point of final analysis

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1.1. The number of minor (e.g. visit time-point violation) or major (e.g. violation of inclusion criteria) protocol deviations collected on a patient participant and site level between randomisation and the defined end of study

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2. Trial withdrawal rate at 6 weeks and 6 months will be measured using the total number of patients randomised in the study. In addition, withdrawal rates will be presented by trial withdrawal reasons and withdrawal stage for both arms.

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3.2. BMI – at randomisation

3.3. Proportion of missing data at the point of final analysis

3.4. Loss to follow up at the point of final analysis

4. Clinical and patient-reported data will be measured by collecting the following information:

4.1. Patient participant-reported outcomes including change in OHS, FJS, WPAI-SHP, SAPS score and quality of life measured using the EQ-5D- 5LTM at 6 weeks and 6 months

4.2. Revision rate, defined as the number of patient participants needing revision arthroplasty within 6 months of surgery

4.3. Infection rate, measured as the rate of re-operation for infection within 6 months of surgery

4.4. Length of stay, measured in hours between patient admission and initial discharge

4.5. Operation time, measured in minutes as the differences between the start and end of the operation

4.6. Incidence of treatment cross over measured as the surgically reported incidence of a patient receiving an intervention other than the one they were allocated through the randomisation system.

4.7. Incidence of intra-operative surgical complications, the proportion of patient participants experiencing greater than grade III severity will be measured and reported according to Clavien Dindo classification system during patient follow-up

4.8. Incidence of postoperative surgical complications, the proportion of patient participants experiencing greater than grade III severity will be measured and reported according to Clavien Dindo classification system during patient follow-up

5. Completeness of the EQ-5D-5LTM and Healthcare Resource Use Survey will be measured by using the following methods:

5.1. Descriptive statistics will be used to assess the frequency of missing data for the EQ-5D-5LTM and Healthcare Resource Use Survey at the point of final analysis.

5.2. Any patterns in missing data will be identified through the use of descriptive statistics at the point of final analysis

Completion date

30/04/2022

Eligibility

Key inclusion criteria

Workstream 1:

1. Age 18 years and above
 2. Undergoing a primary total hip arthroplasty with either a fully cemented or hybrid implant*
 3. Able to give informed consent prior to randomisation
 4. Able to communicate in both written and spoken English
- * All implants and bone cements that are used must be CE marked and used in accordance with their intended use. There is no limitation as to the manufacturer of prostheses used and surgeons will continue to use the implants that they are familiar with. Furthermore, surgeons will be free to use whichever head/socket material and diameter they feel appropriate. The surgical approach, anaesthetic type, rehabilitation after surgery and other concomitant factors will be as per the surgeons' normal care. These factors will be recorded

Workstream 2:

Patient sample:

5. Have been approached to take part in Workstream 1

Surgeon sample:

6. Consultant orthopaedic surgeon at a Workstream 1 site or a potential site for a future full trial

Research nurse sample - Updated 27/07/2021: Healthcare professional sample:

7. Involved in recruiting to Workstream 1 and/or collecting patient self-reported data

Participant type(s)

Mixed

Healthy volunteers allowed

No

Age group

Mixed

Lower age limit

18 years

Upper age limit

110 years

Sex

All

Total final enrolment

0

Key exclusion criteria

Workstream 1:

1. Previous surgery to the hip joint
2. All procedures with an indication other than osteoarthritis
3. Patient requiring complex total hip arthroplasty surgery, specifically augmentation of the acetabulum (eg structural bone graft or metal augment) and/or shortening/de-rotational osteotomy of the femur at the time of surgery
4. Patients requiring bilateral simultaneous total hip arthroplasty
5. Vision impairment that precludes the completion of PROMS questionnaires

Workstream 2:

All participants:

6. Hearing impairment that precludes communication by standard telephone

Date of first enrolment

06/01/2021

Date of final enrolment

31/07/2021

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Wrightington, Wigan and Leigh NHS Foundation Trust

The Elms

Royal Albert Edward

Infirmery

Wigan Lane

Wigan

England

WN1 2NN

Study participating centre

Royal Devon and Exeter NHS Foundation Trust

Royal Devon & Exeter Hospital

Barrack Road

Exeter

England

EX2 5DW

Study participating centre

Northumbria Healthcare NHS Foundation Trust

Rake Lane
North Shields
England
NE29 8NH

Study participating centre**Cambridge University Hospitals NHS Foundation Trust**

Addenbrookes Hospital
Hills Road
Cambridge
England
CB2 0QQ

Study participating centre**University of Manchester**

Division of Psychology and Mental Health
Coupland 1 Building
Oxford Road
Manchester
England
M13 9PL

Sponsor information

Organisation

Wrightington, Wigan and Leigh Teaching Hospitals NHS Foundation Trust

Funder(s)

Funder type

Government

Funder Name

NIHR Central Commissioning Facility (CCF); Grant Codes: PB-PG-0817-20012

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Data sharing statement to be made available at a later date

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
HRA research summary			26/07/2023	No	No
Other publications	A qualitative study of orthopaedic surgeons' perspectives	08/12/2025	10/12/2025	Yes	No
Other unpublished results	version 1.0	10/05/2022	10/05/2022	No	No
Plain English results	version 1.0	10/05/2022	10/05/2022	No	Yes
Protocol file	version v3.0	29/04/2021	27/07/2021	No	No