Rehabilitation after hip resurfacing arthroplasty (RHA)

Submission date	Recruitment status	Prospectively registered
23/04/2010	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
23/04/2010	Completed	[X] Results
Last Edited	Condition category	[] Individual participant data
22/07/2013	Musculoskeletal Diseases	

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

Protocol serial number 4985

Study information

Scientific Title

Evaluation of a specific physiotherapy programme following resurfacing arthroplasty - is it more effective at improving function and muscle strength than standard rehabilitation?

Acronym

Rehab after RHA

Study objectives

To evaluate the effectiveness of a post-operative physiotherapy programme specifically designed to the needs of patients with hip resurfacing arthroplasty (RHA) compared to standard protocols based upon total hip arthroplasty (THA) rehabilitation guidelines.

Hypothesis:

A specific rehabilitation programme following hip resurfacing arthroplasty (RHA) will improve functional outcome assessed at one year post surgery.

Design:

Single blind prospective randomised clinical trial.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Oxfordshire Research Ethics Committee (REC) B approved in August 2006 with an amendment in April 2008 (ref: PB-PG-0407-13216)

Study design

Randomised interventional treatment trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Topic: Musculoskeletal; Subtopic: Musculoskeletal (all Subtopics); Disease: Musculoskeletal

Interventions

A specific rehabilitation programme focussing on range of hip flexion, hip extension strength and single stance higher-demand activities. The content of the rehabilitation programme has been developed following appraisal of the existing literature, focus groups with arthroplasty practitioners and by involvement of a group of patients/service users.

Data will be analysed on an intention to treat basis.

Follow up length: 12 months

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

Oxford Hip Score at 6 and 12 months

Key secondary outcome(s))

Mesaured at 6 weeks, 4 months, 12 months:

- 1. Timed sit-to-stand
- 2. Timed Single leg stand
- 3. Maximal isometric torque strength for flexion

Completion date

31/12/2009

Eligibility

Key inclusion criteria

All patients (either sex, no age limit) who are listed to receive a hip resurfacing arthroplasty will be eligible to take part in the study.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

All

Key exclusion criteria

- 1. Undergoing bilateral arthroplasty
- 2. Minimally invasive surgery
- 3. Further lower limb joint surgery is planned within the next twelve months
- 4. Unable to provide informed consent

Date of first enrolment

01/09/2008

Date of final enrolment

31/12/2009

Locations

Countries of recruitment

United Kingdom

England

Study participating centre Windmill Road

Oxford

Sponsor information

Organisation

Nuffield Orthopaedic Centre NHS Trust (UK)

ROR

https://ror.org/0036ate90

Funder(s)

Funder type

Government

Funder Name

National Institute for Health Research (NIHR) (UK)

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

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type

Details

Results articleresults01/09/2013YesNoParticipant information sheetParticipant information sheet11/11/202511/11/2025NoYes