

Rehabilitation after hip resurfacing arthroplasty (RHA)

Submission date 23/04/2010	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 23/04/2010	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 22/07/2013	Condition category Musculoskeletal Diseases	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
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

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Treatment

Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

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 measure

Oxford Hip Score at 6 and 12 months

Secondary outcome measures

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

Overall study start date

01/09/2008

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

Age group

Not Specified

Sex

Both

Target number of participants

Planned Sample Size: 50

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

England

United Kingdom

Study participating centre

Windmill Road

Oxford

United Kingdom

OX3 7LD

Sponsor information

Organisation

Nuffield Orthopaedic Centre NHS Trust (UK)

Sponsor details

Windmill Rd

Headington

Oxford

England

United Kingdom

OX3 7LD

Sponsor type

Hospital/treatment centre

Website

<http://www.noc.nhs.uk/>

ROR

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

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/09/2013		Yes	No