Randomised trial of hip fractures treated with two different types of hip replacements

Submission date	Recruitment status No longer recruiting	Prospectively registered		
28/09/2007		☐ Protocol		
Registration date 28/09/2007	Overall study status Completed	Statistical analysis plan		
		[X] Results		
Last Edited	Condition category	[] Individual participant data		
21/08/2020	Surgery			

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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0181187886

Study information

Scientific Title

Randomised trial of hip fractures treated with two different types of hip replacements

Study objectives

The aim of this study is to undertake a randomised controlled trial comparing two currently used types of hemiarthroplasties (hip replacements) that are used for the treatment of hip fractures. The aim is to determine if any notable and relevant difference exist between the two designs.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised controlled 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

Hip replacement surgery

Interventions

Hemiarthroplasty method 1 vs hemiarthroplasty method 2

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

Length of time of anaesthesia

Secondary outcome measures

Mortality

Overall study start date

08/11/2006

Completion date

08/11/2014

Eligibility

Key inclusion criteria

- 1. Aged over 49 years old
- 2. Acute admission to Peterborough District Hospital with a hip fracture

Participant type(s)

Patient

Age group

Adult

Sex

Not Specified

Target number of participants

600

Total final enrolment

200

Key exclusion criteria

- 1. Patients being treated conservatively
- 2. Patients having surgery under local anaesthesia
- 3. Patients who have a preference for one form of anaesthesia
- 4. Patients (or their next of kin) unable or unwilling to give written informed consent
- 5. Patients in which either the attending surgeon or anaesthetist feels one method of anaesthesia has preference over the other

Date of first enrolment

08/11/2006

Date of final enrolment

08/11/2014

Locations

Countries of recruitment

England

United Kingdom

Study participating centre Peterborough & Stamford Hospitals NHS Foundation Trust Peterborough United Kingdom PE3 6DA

Sponsor information

Organisation

Record Provided by the NHSTCT Register - 2007 Update - Department of Health

Sponsor details

The Department of Health, Richmond House, 79 Whitehall London United Kingdom SW1A 2NL +44 (0)20 7307 2622 dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type

Government

Funder Name

Peterborough & Stamford Hospitals NHS Foundation Trust (UK)

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 summaryNot provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/06/2012	21/08/2020	Yes	No