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

Submission date	Recruitment status	[_] Prospectively registered
28/09/2007	No longer recruiting	[_] Protocol
Registration date	**Overall study status**	[_] Statistical analysis plan
28/09/2007	Completed	[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
Mr M Parker

Contact details
Peterborough & Stamford Hospitals NHS Foundation Trust
Peterborough District Hospital
Thorpe Road
Peterborough
United Kingdom
PE3 6DA
+44
martyn.parker@pbh-tr.nhs.uk

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 summary

Not 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