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	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

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Additional identifiers

Protocol serial number

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

Study type(s)

Treatment

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(s)

Length of time of anaesthesia

Key secondary outcome(s))

Mortality

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

Healthy volunteers allowed

No

Age group

Adult

Sex

Not Specified

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

United Kingdom

England

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

Funder(s)

Funder type

Government

Funder Name

Peterborough & Stamford Hospitals NHS Foundation Trust (UK)

Results and Publications

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
Participant information shee	Participant information sheet	11/11/2025	11/11/2025 No	Yes