# The Bristol hip view for evaluation of fractured neck of femur

| Submission date              | Recruitment status  No longer recruiting | <ul><li>Prospectively registered</li></ul> |  |
|------------------------------|--|--|--|
| 29/09/2006                   |  | ☐ Protocol                                 |  |
| Registration date 29/09/2006 | Overall study status<br>Completed        | Statistical analysis plan                  |  |
|                              |  | [X] Results                                |  |
| Last Edited                  | Condition category                       | Individual participant data                |  |
| 01/10/2014                   | Injury, Occupational Diseases, Poisoning |  |  |

#### Plain English summary of protocol

Not provided at time of registration

## Contact information

## Type(s)

Scientific

#### Contact name

Dr M Bradley

#### Contact details

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# Additional identifiers

**EudraCT/CTIS** number

**IRAS** number

ClinicalTrials.gov number

Secondary identifying numbers

N0234175548

# Study information

#### Scientific Title

#### **Study objectives**

- 1. Will a modified plain radiograph view (X-ray) help in the diagnosis of fractures to the femoral neck to avoid missed fractures?
- 2. Does the proposed radiographic view statistically increase the visibility of fractured femoral necks when reviewed by orthopaedic surgeons and radiologists?

#### 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)

Diagnostic

#### Participant information sheet

#### Health condition(s) or problem(s) studied

Injury, Occupational Diseases, Poisoning: Factures

#### **Interventions**

Our aims are to see if the modified radiographic view, in addition to the standard series of two radiographs, increases the pick-up rate and visibility of femoral neck fractures.

The standard two radiographs that are already done will act as the control.

#### How subjects are approached:

Patients presenting to the emergency department with a history and examination findings consistent with a fractured upper femur will have the study explained by the examining doctor. The patients will be consented to have an additional radiograph as well as the standard series. The patient will be X-rayed as standard and then the additional radiograph will be performed.

All other treatment will be as standard.

#### Intervention Type

Other

#### **Phase**

Not Applicable

#### Primary outcome measure

The anonymised radiographs will be reviewed by anonymised observers, orthopaedic surgeons, A&E staff and radiologists. The anonymised radiographs will be randomised and any abnormality noted by anonymised observers via a questionnaire. The questionnaire will remain confidential and will not hold any patient identification or personal information. Any further information which is evident from the additional view is to be recorded. Note is made of the confidence of the initial diagnosis, and how the additional radiograph has helped. The proposal has been made after discussion with orthopaedic surgeons, A&E staff and the radiographer-lead to ensure the pathway is feasible and will not add significantly to the workload of the department.

#### Secondary outcome measures

Not provided at time of registration

#### Overall study start date

01/01/2006

#### Completion date

31/12/2006

# **Eligibility**

#### Key inclusion criteria

Patients who present with a history and examination compatible with a fracture of the upper femur

#### Participant type(s)

Patient

#### Age group

Adult

#### Sex

**Not Specified** 

#### Target number of participants

Proposed number of patients: 150

#### Key exclusion criteria

Patients will be excluded if they are unable to understand the study details and hence unable to provide written consent. As the study involves a small additional dose of radiation anybody under 60 years old will be excluded

#### Date of first enrolment

01/01/2006

#### Date of final enrolment

31/12/2006

#### Locations

#### Countries of recruitment

England

**United Kingdom** 

# Study participating centre Southmead Hospital

Bristol United Kingdom BS105NB

# Sponsor information

#### Organisation

Record Provided by the NHSTCT Register - 2006 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**

#### Funder Name

NHS R&D Support Funding (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 | 01/01/2013   |            | Yes            | No              |