ISRCTN71177341 https://doi.org/10.1186/ISRCTN71177341

The role of reinfusion drains in reducing transfusion requirements following primary total hip arthroplasty

Submission date 29/09/2006	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [X] Results [_] Individual participant data
Registration date 29/09/2006	**Overall study status** Completed	
Last Edited 15/11/2011	**Condition category** Surgery	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr Robert Alcock

Contact details
Robert Jones & Agnes Hunt
Orthopaedic Hospital
Oswestry
United Kingdom
SY10 7AG

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0196161708

Study information

Scientific Title
The role of reinfusion drains in reducing transfusion requirements following primary total hip arthroplasty: a randomised controlled trial

Study objectives
Does the use of reinfusion drains offer clinically significant benefits in terms of minimising the need for homologous blood transfusion?

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)
Not specified

Study type(s)
Not Specified

Participant information sheet

Health condition(s) or problem(s) studied
Surgery: Total hip replacement (THR)

Interventions
Re-infusion drains vs no re-infusion drains

Intervention Type
Procedure/Surgery

Phase
Not Specified

Primary outcome measure
Requirement for blood transfusion

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/02/2005

Completion date
31/12/2006

Eligibility

Key inclusion criteria
150 patients having total hip replacement (THR)

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
150

Key exclusion criteria
Patients requiring post operative anti-coagulation

Date of first enrolment
01/02/2005

Date of final enrolment
31/12/2006

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Robert Jones & Agnes Hunt
Oswestry
United Kingdom
SY10 7AG

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
Robert Jones and Agnes Hunt Orthopaedic Hospital (UK), NHS R&D Support Funding

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	01/10/2010		Yes	No