

Drains or no drains following a total knee replacement

Submission date 29/09/2006	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 29/09/2006	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 27/09/2011	Condition category Musculoskeletal Diseases	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Mr Mohamed El Masry

Contact details
Flat 7
22 Park Row
Leeds
United Kingdom
LS1 5HA
+44 07974387455
drmedoelmasry@yahoo.co.uk

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0436165551

Study information

Scientific Title

Study objectives

To evaluate the effectiveness of using a post operative drain following total knee replacement by directly measuring the amount of post operative bleeding using an ultrasound scan.

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

Musculoskeletal Diseases: Total knee replacement (TKR)

Interventions

Drains or no drains

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Post operative haematoma measured at 5 days post operation using an ultrasound scan

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/12/2004

Completion date

04/12/2005

Eligibility

Key inclusion criteria

Patients on the waiting list for total knee replacement (normal population)

Participant type(s)

Patient

Age group

Adult

Sex

Not Specified

Target number of participants

Not provided at time of registration

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/12/2004

Date of final enrolment

04/12/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Flat 7

Leeds

United Kingdom

LS1 5HA

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

Leeds Teaching Hospitals NHS Trust (UK)

Funder Name

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	01/01/2010		Yes	No