

Does continuous epidural infusion as post-operative analgesia alter outcomes in patients undergoing joint arthroplasty?

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 28/11/2014	Condition category Signs and Symptoms	<input type="checkbox"/> Individual participant data <input type="checkbox"/> Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0211117958

Study information

Scientific Title

Does continuous epidural infusion as post-operative analgesia alter outcomes in patients undergoing joint arthroplasty?

Study objectives

Not provided at time of registration

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

Health condition(s) or problem(s) studied

Signs and Symptoms: Pain

Interventions

Group 1 - epidural

Group 2 - patient-controlled analgesia

Intervention Type

Procedure/Surgery

Primary outcome measure

Pain score, side effects, patient satisfaction, demographics, time to discharge, blood loss, deviance from the integrated care pathway, loss of movement, time to mobilisation.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/11/2002

Completion date

31/12/2003

Eligibility

Key inclusion criteria

Adult elective primary artheroplasty patients

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

01/11/2002

Date of final enrolment

31/12/2003

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Royal Surrey County Hospital

Guildford

United Kingdom

GU2 7XX

Sponsor information

Organisation

Department of Health (UK)

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type

Government

Website

<http://www.doh.gov.uk>

Funder(s)**Funder type**

Government

Funder Name

Royal Surrey County Hospital NHS 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