ISRCTN38563644 https://doi.org/10.1186/ISRCTN38563644

A randomised controlled trial investigating the efficacy of a local anaesthetic infusion pump for postoperative analgesia in arthroscopic subarachnoid decompression

Submission date 30/09/2004	**Recruitment status** No longer recruiting	[ ] Prospectively registered [ ] Protocol
Registration date 30/09/2004	**Overall study status** Completed	[ ] Statistical analysis plan [ ] Results
Last Edited 08/04/2014	**Condition category** Signs and Symptoms	[ ] Individual participant data [ ] 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
N0188097521

Study information

Scientific Title

Study objectives
Does the pump reduce oral analgesic requirements and reduce pain scores?

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)
Not Specified

Participant information sheet

Health condition(s) or problem(s) studied
Signs and Symptoms: Pain

Interventions
1. One group will receive local anaesthetic in and around the operated area plus pain management.
2. The other group will have the anaesthetic infusion pump attached.

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
Visual analogue pain scores and oral analgesia intake will be measured every day for seven days following surgery

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/08/2001

Completion date
01/12/2006

Eligibility

Key inclusion criteria
Not provided at time of registration

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
As of June 2004: 25 patients recruited

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/08/2001

Date of final enrolment
01/12/2006

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Orthopaedics
Preston
United Kingdom
PR2 9HT

Sponsor information

Organisation
Department of Health

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type
Government

Funder Name
Lancashire Teaching Hospitals 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