ISRCTN49142784 https://doi.org/10.1186/ISRCTN49142784

A comparison of subcutaneous morphine and diamorphine given via patient-controlled analgesia after hip replacement

Submission date 12/09/2003	**Recruitment status** No longer recruiting	[ ] Prospectively registered [ ] Protocol [ ] Statistical analysis plan [ ] Results [ ] Individual participant data [ ] Record updated in last year
Registration date 12/09/2003	**Overall study status** Completed	
Last Edited 05/12/2014	**Condition category** Signs and Symptoms	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr B Brandner

Contact details
Anaesthetics Department
Middlesex Hospital
Mortimer Street
London
United Kingdom
W1N 8AA
+44 (0)20 7636 8333
abc@email.com

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0263048370

Study information

Scientific Title
A comparison of subcutaneous morphine and diamorphine given via patient-controlled analgesia after hip replacement

Study objectives
Is diamorphine better for the patient in terms of pain scores, side effects and feeling of well being?

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
1. Subcutaneous morphine
2. Diamorphine

Intervention Type
Drug

Phase
Not Applicable

Drug/device/biological/vaccine name(s)
Morphine and diamorphine

Primary outcome measure
Subjective visual analogue scores for pain, nausea, sedation and euphoria.

Secondary outcome measures
Not provided at time of registration

Overall study start date
30/03/1999

Completion date
01/09/2004

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
40 patients from Orthopaedic Surgery

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
30/03/1999

Date of final enrolment
01/09/2004

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

Middlesex Hospital
London
United Kingdom
W1N 8AA

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
Hospital/treatment centre

Funder Name
University College London 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