

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	<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 05/12/2014	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

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