ISRCTN14478115 https://doi.org/10.1186/ISRCTN14478115

Prospective blinded randomised placebo controlled trial investigating whether oxycodone modified release reduces parenteral opioid use following intermediate thoracic surgery

Submission date	Recruitment status	[ ] Prospectively registered
29/07/2004	No longer recruiting	[ ] Protocol
Registration date	Overall study status	[ ] Statistical analysis plan
21/09/2004	Completed	[ ] Results
Last Edited	Condition category	[ ] Individual participant data
07/03/2017	Surgery	[ ] Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
03-274

Study information

Scientific Title
Prospective blinded randomised placebo controlled trial investigating whether oxycodone modified release reduces parenteral opioid use following intermediate thoracic surgery

Acronym
OxyPATS

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)
Not Specified

Participant information sheet

Health condition(s) or problem(s) studied
Post-operative pain following intermediate thoracic surgery

Interventions
Oxycodone modified release 10 mg bd or placebo for 48 hrs postoperatively.

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)
Oxycodone

Primary outcome measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/01/2004

Completion date
31/12/2005

Eligibility

Key inclusion criteria
Adult patients scheduled for unilateral intermediate thoracic surgery (e.g. pleural abrasion, talc pleurodesis and lung biopsy)

Participant type(s)
Patient

Age group
Not Specified

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
01/01/2004

Date of final enrolment
31/12/2005

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Department of Anaesthesia
Harefield
United Kingdom
UB9 6JH

Sponsor information

Organisation
Harefield Hospital (UK)

Sponsor details
Dr CPR Walker
Department of Anaesthesia & Pain Management
Hill End Road
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Sponsor type
Hospital/treatment centre

ROR
https://ror.org/04fwa4t58

Funder(s)

Funder type
Industry

Funder Name
Royal Brompton & Harefield NHS Trust fund the salaries.

Funder Name
Napp Pharmaceuticals are supplying the study drugs and placebos.

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