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

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr David Royston

Contact details
Department of Anaesthesia
Harefield Hospital
Hill End Road
Harefield
United Kingdom
UB9 6JH

Additional identifiers

Protocol serial number
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

Study type(s)
Not Specified

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(s)
Not provided at time of registration

Key secondary outcome(s))
Not provided at time of registration

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

Healthy volunteers allowed
No

Age group
Not Specified

Sex
Not Specified

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
United Kingdom

England

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

Sponsor information

Organisation
Harefield Hospital (UK)

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

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration