

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

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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