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

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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
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# Sponsor information

**Organisation**
Harefield Hospital (UK)

**Sponsor details**
Dr CPR Walker
Department of Anaesthesia & Pain Management
Hill End Road
Harefield
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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