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Double blind comparison of analgesia with morphine or morphine and ketamine after abdominal surgery in children aged 3 months to 1 year

Submission date 12/09/2003	**Recruitment status** No longer recruiting	[ ] Prospectively registered [ ] Protocol
Registration date 12/09/2003	**Overall study status** Completed	[ ] Statistical analysis plan [ ] Results
Last Edited 02/02/2017	**Condition category** Signs and Symptoms	[ ] 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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0012120218

Study information

Scientific Title
Double blind comparison of analgesia with morphine or morphine and ketamine after abdominal surgery in children aged 3 months to 1 year

Study objectives
Does the use of ketamine with morphine for acute postoperative pain reduce inflammatory hypersensitivity and morphine requirements?

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
Not available in web format, please use contact details to request a participant information sheet

Health condition(s) or problem(s) studied
Signs and Symptoms: Pain

Interventions
1. Post-operative pain therapy
2. Standard care

Intervention Type
Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)
Ketamine, morphine

Primary outcome measure
Evaluation of analgesic technique

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/09/2002

Completion date
16/04/2007

Eligibility

Key inclusion criteria
Children who have undergone abdominal surgery

Participant type(s)
Patient

Age group
Child

Sex
Both

Target number of participants
Not provided at time of registration

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/09/2002

Date of final enrolment
16/04/2007

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Great Ormond Street Hospital
London
United Kingdom
WC1N 3JH

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
Government

Funder Name
Great Ormond Street Hospital for Children NHS Trust / Institute of Child Health (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