ISRCTN69774569 https://doi.org/10.1186/ISRCTN69774569

Comparison of caudal epidural using 0.25% bupivacaine and 0.5 mg/kg ketamine with penile block using 0.5% bupivacaine for postoperative analgesia following day case circumcision.

Submission date 30/09/2004	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [X] Results [_] Individual participant data
Registration date 30/09/2004	**Overall study status** Completed	
Last Edited 15/07/2009	**Condition category** Surgery	

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
N0185139342

Study information

Scientific Title

Study objectives
The null hypothesis is that there is no difference in the quality and duration of post-operative analgesia after day-case circumcisions afforded by a caudal block using a mixture of bupivacaine and ketamine and a penile block using bupivacaine.

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)
Other

Participant information sheet

Health condition(s) or problem(s) studied
Surgery: Analgesia

Interventions
Double blinded randomised controlled trial

Intervention Type
Procedure/Surgery

Phase
Not Specified

Primary outcome measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/09/2002

Completion date
31/03/2005

Eligibility

Key inclusion criteria
44 subjects all aged less than 16 years having elective day-case circumcisions.

Participant type(s)
Patient

Age group
Child

Upper age limit
16 Years

Sex
Male

Target number of participants
44

Key exclusion criteria
Does not match inclusion criteria

Date of first enrolment
01/09/2002

Date of final enrolment
31/03/2005

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Anaesthetic Department
Plymouth
United Kingdom
PL6 8DH

Sponsor information

Organisation
Department of Health

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
Plymouth Hospitals NHS Trust (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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/12/2008		Yes	No