

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
Registration date 30/09/2004	Overall study status Completed	☐ Statistical analysis plan ☒ Results
Last Edited 15/07/2009	Condition category Surgery	☐ Individual participant data

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
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

Primary study design

Interventional

Study design

Randomised controlled trial

Study type(s)

Other

Health condition(s) or problem(s) studied

Surgery: Analgesia

Interventions

Double blinded randomised controlled trial

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome(s)

Not provided at time of registration

Key secondary outcome(s)

Not provided at time of registration

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

Healthy volunteers allowed

No

Age group

Child

Upper age limit

16 Years

Sex

Male

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

United Kingdom

England

Study participating centre

Anaesthetic Department

Plymouth

United Kingdom

PL6 8DH

Sponsor information

Organisation

Department of Health

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Plymouth Hospitals NHS Trust (UK)

Results and Publications

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