

# Does the route of administration of ketamine influence perioperative and postoperative analgesia?

<b>Submission date</b> 12/09/2003	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 12/09/2003	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
<b>Last Edited</b> 16/09/2014	<b>Condition category</b> Signs and Symptoms	<input type="checkbox"/> Individual participant data <input type="checkbox"/> Record updated in last year

## Plain English summary of protocol

Not provided at time of registration

## Contact information

### Type(s)

Scientific

### Contact name

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### Contact details

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## Additional identifiers

### Protocol serial number

N0084113941

## Study information

Scientific Title

**Study objectives**

To compare the duration of analgesia with ketamine given by two different routes.

**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)**

Treatment

**Health condition(s) or problem(s) studied**

Signs and Symptoms: Pain

**Interventions**

1. Caudal ketamine
2. Intravenous ketamine
3. No ketamine

**Intervention Type**

Other

**Phase**

Not Applicable

**Primary outcome(s)**

1. Duration of postoperative analgesia
2. Nausea, sedation and motor blockade

**Key secondary outcome(s)**

Not provided at time of registration

**Completion date**

01/06/2004

**Eligibility****Key inclusion criteria**

Children undergoing herniotomy, orchidoplexy and repair of hydrocoele

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Child

**Sex**

Not Specified

**Key exclusion criteria**

Not provided at time of registration

**Date of first enrolment**

28/06/2002

**Date of final enrolment**

01/06/2004

**Locations****Countries of recruitment**

United Kingdom

England

**Study participating centre**

Department of Anaesthetics

Hull

United Kingdom

HU3 2JZ

**Sponsor information****Organisation**

Department of Health (UK)

**Funder(s)****Funder type**

Government

**Funder Name**

The North and South Bank Research and Development Consortium (UK)

## **Results and Publications**

**Individual participant data (IPD) sharing plan**

**IPD sharing plan summary**

Not provided at time of registration