ISRCTN86456561 https://doi.org/10.1186/ISRCTN86456561

Intraperitoneal bupivacaine for post-operative analgesia

Submission date	Recruitment status	[ ] Prospectively registered
30/09/2004	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
30/09/2004	Completed	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
10/11/2022	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

Protocol serial number
N0256130924

Study information

Scientific Title
Intraperitoneal bupivacaine for post-operative analgesia

Study objectives

Does Adept prolong the action of local anaesthetic?

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)
Not Specified

Health condition(s) or problem(s) studied
Signs and Symptoms: Post-operative pain

Interventions
Randomised controlled trial
Group 1: Bupivacaine + Saline (control)
Group 2: Bupivacaine + Adept

Intervention Type
Drug

Phase
Not Applicable

Drug/device/biological/vaccine name(s)
Bupivacaine

Primary outcome(s)
Service outcome development

Key secondary outcome(s)
Not provided at time of registration

Completion date
31/08/2004

Eligibility

Key inclusion criteria
35 patients in each group

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Adult

Sex
Female

Key exclusion criteria
Does not match inclusion criteria

Date of first enrolment
04/11/2003

Date of final enrolment
31/08/2004

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
University Department of Obstetrics and Gynaecology
London
United Kingdom
NW3 2QG

Sponsor information

Organisation
Department of Health

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
The Royal Free Hampstead NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan
Not provided at time of registration

IPD sharing plan summary
Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Other publications	feasibility study	01/06/2004		Yes	No