

Post-operative pain relief after vitreo-retinal surgery with Subtenon L Bupivacaine infiltration

Submission date 30/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 19/09/2017	Condition category Surgery	<input type="checkbox"/> 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
N0064131841

Study information

Scientific Title
Post-operative pain relief after vitreo-retinal surgery with Subtenon L Bupivacaine infiltration: a randomised controlled trial

Study objectives

Does Subtenon L Bupivacaine significantly reduce requirements for post-operative analgesia after vitreo retinal surgery compared with current analgesia?

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

Surgery: Analgesia

Interventions

Subtenon L Bupivacaine vs current analgesia

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

Subtenon L Bupivacaine

Primary outcome(s)

Not provided at time of registration

Key secondary outcome(s)

Not provided at time of registration

Completion date

01/04/2004

Eligibility**Key inclusion criteria**

Not provided at time of registration

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/11/2003

Date of final enrolment

01/04/2004

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

Academic Unit of Ophthalmology

Birmingham

United Kingdom

B18 7HQ

Sponsor information**Organisation**

Department of Health

Funder(s)**Funder type**

Government

Funder Name

Sandwell and West Birmingham 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	01/02/2005		Yes	No