

Assessment of local anaesthetic techniques in cataract surgery.

Submission date 30/09/2005	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 30/09/2005	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 11/02/2008	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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0065149380

Study information

Scientific Title

Study objectives

To evaluate the pain patients experience during cataract surgery with three different types of local anaesthesia, using two types of pain questionnaires (McGill pain questionnaire and visual analogue scale).

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)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Surgery: Cataract

Interventions

A randomised, controlled trial comparing three types of anaesthesia: peribulbar, sub-tenon and topical.

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

Quality of pain control through cataract surgery with different types of anaesthetic. Assessment of pain the patients have with anaesthetic administration.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/10/2004

Completion date

01/10/2006

Eligibility

Key inclusion criteria

Not provided at time of registration

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

100 patients in each arm - 300

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/10/2004

Date of final enrolment

01/10/2006

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Department of Ophthalmology

Sunderland

United Kingdom

SR2 9HP

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL
+44 (0)20 7307 2622
dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

<http://www.dh.gov.uk/Home/fs/en>

Funder(s)

Funder type

Government

Funder Name

City Hospitals Sunderland NHS Trust R&D Department

Funder Name

NHS R&D Support Funding

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?
Abstract results	publication in Br J Ophthalmol. Sep;89(9):1228 entitled "Patient satisfaction with anaesthesia comparing sub-Tenon's block and topical anaesthesia" but no abstract available	01/09/2005		No	No