

A comparison of peribulbar and sub-Tenon's techniques for bilateral cataract surgery

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|--|---|---|
| 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 21/01/2009 | 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
N0282136776

Study information

Scientific Title

Study objectives

To compare peribulbar and sub-Tenon's local anaesthetics in patients having cataract surgery on both eyes

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)

Not specified

Study type(s)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Surgery: Cataract

Interventions

Peribulbar vs sub-Tenon's local anaesthetics

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

Information on most appropriate technique

Secondary outcome measures

Not provided at time of registration

Overall study start date

30/01/2004

Completion date

30/01/2005

Eligibility

Key inclusion criteria

100 patients/200 procedures.

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

100

Key exclusion criteria

1. Children
2. Surgery other than cataracts
3. Patients with axial length greater than 27 mm
4. Patients with only one eye

Date of first enrolment

30/01/2004

Date of final enrolment

30/01/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Worcestershire Royal Hospital

Worcester

United Kingdom

WR5 1DD

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type

Government

Website

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

Funder(s)**Funder type**

Government

Funder Name

Worcestershire Acute Hospitals NHS Trust (UK)

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? |
|---------------------------------|---------|--------------|------------|----------------|-----------------|
| Results article | results | 01/01/2009 | | Yes | No |