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Sensory nerve block for upper lid surgery: a prospective randomised comparative study

Submission date
29/09/2006

Recruitment status
Stopped

Registration date
29/09/2006

Overall study status
Stopped

Last Edited
27/09/2011

Condition category
Surgery

[ ] Prospectively registered
[ ] Protocol
[ ] Statistical analysis plan
[ ] Results
[ ] Individual participant data
[ ] 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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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0214163271

Study information

Scientific Title

Study objectives
To evaluate a combination regional block of the frontal and lacrimal nerves by the Hildreth-Silver technique for upper lid surgery.

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: Eye

Interventions
Hildreth-Silver technique vs other technique

Added 09 September 2008: trial stopped due to unsatisfactory results.

Intervention Type
Procedure/Surgery

Phase
Not Specified

Primary outcome measure
Pain scores for injection and procedures, complications of anaesthesia, need for supplementary anaesthesia, levator function, patient co-operation score, surgical features, anaesthetic records

Secondary outcome measures
Not provided at time of registration

Overall study start date
21/01/2005

Completion date
31/12/2005

Reason abandoned (if study stopped)
Unsatisfactory results

Eligibility

Key inclusion criteria
Any upper lid surgery suitable for local anaesthesia

Participant type(s)
Patient

Age group
Adult

Sex
Not Specified

Target number of participants
40

Key exclusion criteria
1. Age<16 years
2. Previous lid surgery
3. Difficulty in comprehension of study method or processes.

Date of first enrolment
21/01/2005

Date of final enrolment
31/12/2005

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

The Wolverhampton & Midland Counties Eye Infirmary
Wolverhampton
United Kingdom
WV3 9QR

Sponsor information

Organisation
Record Provided by the NHSTCT Register - 2006 Update - Department of Health

Sponsor details
The Department of Health, 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
The Royal Wolverhampton Hospitals NHS Trust (UK), 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