

Bleb Dysesthesia Trial

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 type="checkbox"/> Results
Last Edited 29/05/2014	Condition category Surgery	<input type="checkbox"/> Individual participant data <input type="checkbox"/> 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

N0203125674; N0199129912; N0176115690

Study information

Scientific Title

A randomised trial of the effect of bleb site on eye comfort following trabeculectomy

Study objectives

To investigate long-term post-operative eye discomfort following glaucoma surgery; in particular whether two different wound locations and therefore eye fluid drainage, differ in the level of eye comfort experienced as measured by a validated questionnaire, the 'Glaucoma Symptom Scale'.

On 14/07/2011 the following changes were made to this trial record:

1. The end date was changed from 31/12/2008 to 30/06/2010.
2. The target number of participants was changed from 40 to 150.
3. The trial was originally a multi-centre trial but due to recruitment issues at other centres the Oxford site proceeded single-handedly.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Oxford Research Ethics Committee B, 10/09/2002, ref: C02.200

Study design

Randomised controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Other

Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Surgery: Trabeculectomy

Interventions

The study design is a randomised trial designed to compare two accepted variations in surgical techniques of trabeculectomy and its effect on postoperative eye comfort. Patients who are undergoing first trabeculectomy will be approached to participate in the study. Patients will either be recruited in person or by invitation letter. When the patient gives written consent, they are randomised either to superior trabeculectomy or superonasal trabeculectomy. At the pre-operative visit they will answer a 10-question self-administered questionnaire, the Glaucoma Symptom Scale. Some pre-operative data will be collected on study forms identified only by hospital number and study number.

During the trabeculectomy, the bleb will be placed below the middle upper lid, or placed below the more inner upper lid.

After the trabeculectomy, usual medical care will take place uninfluenced by their participation in this study. At the 6-month post-operative visit, the patient will be examined for the study and a second Glaucoma Symptom Scale questionnaire will be completed by the patient. The examination will record clinical findings such as visual acuity, eye pressure and eye examination findings. The study ends here and patients continue their usual care.

Intervention Type

Procedure/Surgery

Phase

Not Applicable

Primary outcome measure

1. Eye comfort at 6 months post-surgery as measured by the Glaucoma Symptom Scale
2. Eye features associated with eye discomfort (dysesthesia) at 6 months post-surgery

Secondary outcome measures

No secondary outcome measures

Overall study start date

10/09/2002

Completion date

30/06/2010

Eligibility

Key inclusion criteria

1. Aged 18 years and over, either sex
2. Listed for trabeculectomy
3. Satisfy the inclusion criteria

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

150

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

10/09/2002

Date of final enrolment

30/06/2010

Locations**Countries of recruitment**

England

United Kingdom

Study participating centre

John Radcliffe Hospital

Oxford

United Kingdom

OX3 9DU

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

Hospital/treatment centre

Funder Name

Royal Devon and Exeter NHS Trust (UK)

Funder Name

Royal Berkshire and Battle Hospitals NHS Trust (UK)

Funder Name

Oxford Radcliffe 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