Prospective comparison of standard LASIK (= laser assisted in situ keratomileusis) vs cross cylinder ablation LASIK for the correction of post keratoplasty ametropias

Submission date	Recruitment status	Prospectively registered
30/09/2005	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
30/09/2005	Completed	Results
Last Edited	Condition category	Individual participant data
16/03/2016	Eye Diseases	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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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0141156870

Study information

Scientific Title

Prospective comparison of standard LASIK (= laser assisted in situ keratomileusis) vs cross cylinder ablation LASIK for the correction of post keratoplasty ametropias

Study objectives

To compare 2 different methods of applying the excimer laser with LASIK to correct refractive errors following full thickness corneal graft 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)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Eye Diseases: Ametropia

Interventions

Prospective double masked randomised study

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Reduction in corneal astigmatism

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/03/2005

Completion date

01/04/2006

Eligibility

Key inclusion criteria

LASIK patients

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

128

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/03/2005

Date of final enrolment

01/04/2006

Locations

Countries of recruitment

England

United Kingdom

Study participating centre Moorfields Eye Hospital

London United Kingdom EC1V 2PD

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

Moorfields Eye Hospital NHS Foundation 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