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

Submission date 30/09/2005	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol
Registration date 30/09/2005	**Overall study status** Completed	[_] Statistical analysis plan [_] Results
Last Edited 16/03/2016	**Condition category** Eye Diseases	[_] Individual participant data [_] Record updated in last year

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
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
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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