

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	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 30/09/2005	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 16/03/2016	Condition category Eye Diseases	<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

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Additional identifiers

Protocol serial number

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

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Eye Diseases: Ametropia

Interventions

Prospective double masked randomised study

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Reduction in corneal astigmatism

Key secondary outcome(s))

Not provided at time of registration

Completion date

01/04/2006

Eligibility**Key inclusion criteria**

LASIK patients

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

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

United Kingdom

England

Study participating centre

Moorfields Eye Hospital

London

United Kingdom

EC1V 2PD

Sponsor information

Organisation

Department of Health

Funder(s)

Funder type

Government

Funder Name

Moorfields Eye Hospital NHS Foundation Trust (UK) NHS R&D Support Funding

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration