

Corneal protection after cataract surgery

Submission date 28/09/2007	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 28/09/2007	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 12/05/2017	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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0214181354

Study information

Scientific Title

Corneal protection after cataract surgery

Study objectives

What is the most comfortable and safest method for corneal protection after cataract 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)

Hospital

Study type(s)

Treatment

Participant information sheet

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

Health condition(s) or problem(s) studied

Eye diseases: cataract

Interventions

Eyepad vs eyeshield vs viscoelastic agent plus eyeshield in patients who have undergone cataract surgery.

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

1. Pain scores on first postoperative day
2. Corneal fluorescein staining on first postoperative day

Secondary outcome measures

Patient mobility and satisfaction in the post-operative period

Overall study start date

07/04/2006

Completion date

07/06/2006

Eligibility

Key inclusion criteria

Patients who have undergone eye surgery, undergoing uncomplicated phacoemulsification under local anaesthesia

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

120

Key exclusion criteria

1. Age < 18 years old
2. Difficulty in comprehension of study method or processes
3. Complicated cataract surgery
4. Pre-existing corneal disease
5. Corneal wound requiring suturing
6. Postoperative intraocular pressure of >25 mmHg
7. Greater than moderate anterior chamber reaction (+2 cells/flare) postoperatively

Date of first enrolment

07/04/2006

Date of final enrolment

07/06/2006

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Wolverhampton Eye Infirmary
Wolverhampton
United Kingdom
WV10 0QP

Sponsor information

Organisation

Record Provided by the NHSTCT Register - 2007 Update - Department of Health

Sponsor details

The Department of Health
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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), RWHT in kind contribution, 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