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Corneal protection after cataract surgery

Submission date	Recruitment status	[ ] Prospectively registered
28/09/2007	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
28/09/2007	Completed	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
12/05/2017	Eye Diseases	[ ] 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
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

Primary study design
Interventional

Study design
Randomised controlled trial

Study type(s)
Treatment

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

Key secondary outcome(s)
Patient mobility and satisfaction in the post-operative period

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

Healthy volunteers allowed
No

Age group
Adult

Sex
All

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
United Kingdom

England

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

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

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration