

Corneal protection after cataract surgery

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

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