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Corneal protection after cataract surgery

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

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