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Preferred treatment for removal of corneal rust

Submission date 30/09/2005	**Recruitment status** No longer recruiting	[ ] Prospectively registered [ ] Protocol [ ] Statistical analysis plan [ ] Results [ ] Individual participant data [ ] Record updated in last year
Registration date 30/09/2005	**Overall study status** Completed	
Last Edited 11/04/2014	**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

Protocol serial number
N0185146360

Study information

Scientific Title

Study objectives
To identify what is best practice for removal of corneal rust, ensuring that patient care is not compromised.

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)
Not Specified

Health condition(s) or problem(s) studied
Eye Diseases: Corneal damage

Interventions
Patients will be randomly allocated 2 or 5 days using sealed envelopes.
The following will be undertaken: 1.Case History, 2.Visual Acuity, 3. Examination: Slit lamp bio microscopy 4. Check whether patient fits criteria, 5. ask patient to participate, give patient information sheet, obtain consent. 6. Nurse practitioner opens envelope to identify 2 or 5 days. 7. Size and depth of foreign body will be measured and documented. 8. Other signs, symptoms documented eg inflammation. 9. pain score 1-10 analogue scale 10. procedure of removal 11. measurement of rust remaining 12 treatment given i.e medication, whether padded 13, follow up appointment according to random allocation 14 repeat visits to follow above procedure 15 telephone call to be made if follow up appointment not attended.

Intervention Type
Other

Phase
Not Specified

Primary outcome(s)
Aims & Objectives: To identify what is the best practice for removal of cornea rust, ensuring that patient care is not compromised. To compare differences in discomfort felt by the patient and healing process of the cornea when visits are at 2 or 5 day intervals. Study endpoints: To compare ease and completion of removal of cornea rust.

Key secondary outcome(s)
Not provided at time of registration

Completion date
28/02/2004

Eligibility

Key inclusion criteria

Not provided at time of registration

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
27/02/2003

Date of final enrolment
28/02/2004

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
Accident & Emergency OPD Department
Plymouth
United Kingdom
PL4 6PL

Sponsor information

Organisation
Department of Health

Funder(s)

Funder type
Government

Funder Name
Plymouth Hospitals NHS Trust (UK), Own Account

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration