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Do cycloplegics lessen the pain for a patient in the first 24 h following a corneal abrasion?

Submission date	Recruitment status	[ ] Prospectively registered
12/09/2003	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
12/09/2003	Completed	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
07/12/2015	Injury, Occupational Diseases, Poisoning	[ ] 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
N0557119809

Study information

Scientific Title
Do cycloplegics lessen the pain for a patient in the first 24 h following a corneal abrasion?

Study objectives
Does the application of a topical cycloplegic have an effect upon the pain experienced by patients in the 24 h following a simple traumatic corneal abrasion?

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)
Not specified

Study type(s)
Treatment

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

Health condition(s) or problem(s) studied
Signs and Symptoms: Pain

Interventions
Prospective, double blinded, randomized controlled trial. Random allocation to:
1. Cyclopentolate
2. Normal saline

Intervention Type
Drug

Phase
Not Applicable

Drug/device/biological/vaccine name(s)
Cyclopentolate

Primary outcome measure

Pain over the subsequent 24 h

Secondary outcome measures
Not provided at time of registration

Overall study start date
14/01/2003

Completion date
31/07/2003

Eligibility

Key inclusion criteria
100 patients

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
100

Key exclusion criteria
Does not match inclusion criteria

Date of first enrolment
14/01/2003

Date of final enrolment
31/07/2003

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

Russells Hall Hospital
Dudley
United Kingdom
DY1 2HQ

Sponsor information

Organisation
Department of Health (UK)

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
http://www.doh.gov.uk

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
The Dudley Group of Hospitals NHS Trust (UK)

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