

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

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 07/12/2015	Condition category Injury, Occupational Diseases, Poisoning	<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
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

Study type(s)

Treatment

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(s)

Pain over the subsequent 24 h

Key secondary outcome(s))

Not provided at time of registration

Completion date

31/07/2003

Eligibility

Key inclusion criteria

100 patients

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

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**

United Kingdom

England

Study participating centre

Russells Hall Hospital

Dudley

United Kingdom

DY1 2HQ

Sponsor information**Organisation**

Department of Health (UK)

Funder(s)**Funder type**

Hospital/treatment centre

Funder Name

The Dudley Group of Hospitals NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration