

Randomised controlled study comparing the effectiveness of Geliperm and Lacrilube in the prevention of corneal damage in the critically ill.

Submission date 30/09/2005	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 30/09/2005	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
Last Edited 28/06/2010	Condition category Eye Diseases	<input type="checkbox"/> Individual participant data

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

N0205151291

Study information

Scientific Title

Study objectives

This project has a simple objective: to evaluate the effectiveness of Geliperm and Lacrilube at preventing corneal damage.

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

Health condition(s) or problem(s) studied

Eye Diseases: Corneal damage

Interventions

Ocular lubricant (Lacrilube) vs polyacrylamide hydrogel dressings (Geliperm)

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

31/08/2004

Completion date

01/03/2005

Eligibility

Key inclusion criteria

All patients admitted to ITU will be considered for recruitment.

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

Added 28/06/10: 80 eyes from 40 patients

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

31/08/2004

Date of final enrolment

01/03/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

ITU

London

United Kingdom

E1 1BB

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL
+44 (0)20 7307 2622
dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

<http://www.dh.gov.uk/Home/fs/en>

Funder(s)

Funder type

Government

Funder Name

Barts and The London NHS Trust (UK) 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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
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[Results article](#)

results

01/03/2009

Yes

No