ISRCTN60137486 https://doi.org/10.1186/ISRCTN60137486

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	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [X] Results [_] Individual participant data
Registration date 30/09/2005	**Overall study status** Completed	
Last Edited 28/06/2010	**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

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
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United Kingdom
SW1A 2NL
+44 (0)20 7307 2622
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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?

Results article	results	01/03/2009	Yes	No