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A randomised prospective study to evaluate blepharoplasty skin closure by Tisseel® glue in comparison with conventional suturing technique

Submission date
29/09/2006

Recruitment status
No longer recruiting

Registration date
29/09/2006

Overall study status
Completed

Last Edited
02/06/2017

Condition category
Surgery

[_] Prospectively registered
[_] Protocol
[_] Statistical analysis plan
[_] Results
[_] Individual participant data
[_] 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
N0016173487

Study information

Scientific Title
A randomised prospective study to evaluate blepharoplasty skin closure by Tisseel® glue in comparison with conventional suturing technique

Study objectives
What is the wound characteristics of glue closure in the eyelid?

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
Suturing techniques

Interventions
Patients were randomised between:
1. Tisseel® glue
2. Conventional suturing technique

Intervention Type
Drug

Phase
Not Applicable

Drug/device/biological/vaccine name(s)
Tisseel® glue

Primary outcome measure
Independent observer wounds and questionnaire patient

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/01/2004

Completion date
01/12/2005

Eligibility

Key inclusion criteria
Not provided at time of registration

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
50

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/01/2004

Date of final enrolment
01/12/2005

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Ophthalmological Surgery
London
United Kingdom
W6 8RF

Sponsor information

Organisation
Record Provided by the NHSTCT Register - 2006 Update - Department of Health (UK)

Sponsor details
The Department of Health
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
Hammersmith Hospital NHS Trust (UK)

Funder Name
NHS R&D Support Funding 2004/05 (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