

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

<b>Submission date</b> 29/09/2006	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
<b>Registration date</b> 29/09/2006	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
<b>Last Edited</b> 02/06/2017	<b>Condition category</b> Surgery	<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

### Contact name

Miss Jane Olver

### Contact details

Ophthalmological Surgery  
Charing Cross Hospital  
Fulham Palace Road  
Hammersmith  
London  
United Kingdom  
W6 8RF  
+44 (0)20 8846 1497  
j.olver@imperial.ac.uk

## 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