

# The role of bioglue in eliminating prolonged alveolar air leak in thoracic surgery

<b>Submission date</b> 12/09/2003	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 12/09/2003	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
<b>Last Edited</b> 20/07/2007	<b>Condition category</b> Surgery	<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**  
N0201118681

# Study information

## Scientific Title

## Study objectives

Will bioglue reduce required length of chest drainage and hospital stay by reducing persistent alveolar air leaks following major lung resection?

## 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)

Not Specified

## Participant information sheet

## Health condition(s) or problem(s) studied

Surgery: Thoracic

## Interventions

Surgical treatment only vs surgical treatment and BioGlue

## Intervention Type

Procedure/Surgery

## Phase

Not Specified

## Primary outcome measure

1. Duration of air leak
2. Duration of intercostal drainage
3. Duration of hospital stay
4. Incidence of complications

**Secondary outcome measures**

Not provided at time of registration

**Overall study start date**

01/07/2002

**Completion date**

31/01/2005

## Eligibility

**Key inclusion criteria**

Not provided at time of registration

**Participant type(s)**

Patient

**Age group**

Not Specified

**Sex**

Both

**Target number of participants**

104 patients

**Key exclusion criteria**

Not provided at time of registration

**Date of first enrolment**

01/07/2002

**Date of final enrolment**

31/01/2005

## Locations

**Countries of recruitment**

England

United Kingdom

**Study participating centre**

Surgery

London

United Kingdom

SW3 6NP

# Sponsor information

## Organisation

Department of Health (UK)

## Sponsor details

Richmond House

79 Whitehall

London

United Kingdom

SW1A 2NL

## Sponsor type

Government

## Website

<http://www.doh.gov.uk>

# Funder(s)

## Funder type

Government

## Funder Name

Royal Brompton and Harefield NHS Trust (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

## Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Results article</a>	Results:	01/07/2006		Yes	No

