

Comparison of urokinase versus primary video-assisted thorascopic surgery for the treatment of empyema in children

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 15/05/2009	Condition category Respiratory	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr A Jaffe

Contact details

Portex Unit
Institute of Child Health
30 Guilford Street
London
United Kingdom
WC1N 1EH
+44 (0)20 7405 9200

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0012114253

Study information

Scientific Title

Study objectives

Is primary video-assisted thorascopic surgery (VATS) the treatment of choice for empyema in children?

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

Respiratory: Empyema

Interventions

Random allocation to (A) Treatment for empyema (B) Placebo.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Number of hospital days post procedure until ready for discharge.

Secondary outcome measures

1. Total length of hospital stay
2. Total length of illness
3. Number of chest tube days

Overall study start date

01/02/2002

Completion date

31/01/2005

Eligibility

Key inclusion criteria

Children with empyema.

Participant type(s)

Patient

Age group

Child

Sex

Not Specified

Target number of participants

Not provided at time of registration

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/02/2002

Date of final enrolment

31/01/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Portex Unit
London
United Kingdom
WC1N 1EH

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
Great Ormond Street Hospital for Children NHS Trust / Institute of Child Health (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?
Results article	results	15/07/2006		Yes	No