

Tidal Volume - Cytokine levels trial in anaesthetised patients undergoing positive pressure ventilation

Submission date 30/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 26/09/2016	Condition category 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
Dr Bernadette Ewah

Contact details
Epsom and St Helier NHS Trust
Epsom General Hospital
Dorking Road
Epsom
Surrey
United Kingdom
KT18 7EG
+44 (0)1372 7355270
bewah99@yahoo.com

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0112112997

Study information

Scientific Title

Tidal Volume - Cytokine levels trial in anaesthetised patients undergoing positive pressure ventilation

Study objectives

What tidal volume is beneficial for artificial ventilation during surgery?

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)

Not specified

Study type(s)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Surgery: Ventilation

Interventions

Patients randomly divided into two groups. One group is ventilated with a standard tidal volume of 10 ml/kg. The second group is ventilated with 6 ml/kg. Cytokine levels in bronchial aspirate and blood are measured.

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

Cytokine levels.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/02/2002

Completion date

01/02/2004

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

01/02/2002

Date of final enrolment

01/02/2004

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Epsom and St Helier NHS Trust
Surrey
United Kingdom
KT18 7EG

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type

Government

Website

<http://www.dh.gov.uk/Home/fs/en>

Funder(s)

Funder type

Government

Funder Name

Epsom and St Helier University Hospitals 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