

Randomised controlled trial of closed circuit Continuous Positive Airway Pressure (CPAP) and open circuit CPAP with a Boussignac valve for the treatment of acute pulmonary oedema.

Submission date 30/09/2004	Recruitment status No longer recruiting	☐ Prospectively registered ☐ Protocol
Registration date 30/09/2004	Overall study status Completed	☐ Statistical analysis plan ☒ Results
Last Edited 10/07/2009	Condition category Respiratory	☐ 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

Protocol serial number
N0013101032

Study information

Scientific Title

Study objectives

This study will seek to determine whether the Boussignac valve system is as effective as conventional CPAP using a Drager CF800 circuit.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Primary study design

Interventional

Study design

Randomised controlled trial

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Respiratory: Acute pulmonary oedema

Interventions

The patient will be randomised to one of two groups:

1. CPAP with the closed circuit Drager CF800 system

OR

2. CPAP with the open circuit Vygon Boussignac valve system

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

The main outcome measure will be the PaCO₂ at 30 and 60 minutes. Other outcome measures will be PaO₂ at 30 and 60 minutes, need for intubation, and tolerance of mask/circuit.

Key secondary outcome(s)

Not provided at time of registration

Completion date

31/12/2003

Eligibility**Key inclusion criteria**

Patients with acute pulmonary oedema attending Accident and Emergency (A&E). We intend to recruit 50 patients to allow for recruitment errors and drop outs etc.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

Does not meet inclusion criteria

Date of first enrolment

01/10/2001

Date of final enrolment

31/12/2003

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

Acute Medicine

London

United Kingdom

SE1 7EH

Sponsor information**Organisation**

Department of Health

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Guy's and St Thomas' NHS Trust (UK)

Results and Publications

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	01/06/2005		Yes	No