

A randomised trial of elective continuous positive airway pressure (CPAP) verses rescue CPAP after extubation in infants following cardiac surgery

Submission date 29/09/2006	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 29/09/2006	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 18/05/2017	Condition category Neonatal Diseases	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<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 Ruth Helen Wakeman

Contact details

Royal Brompton & Harefield NHS Trust
Sydney Street
London
United Kingdom
SW3 6NP
+44 (0)20 7351 8088/8436
r.wakeman@rbht.nhs.uk

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0201170878

Study information

Scientific Title

A randomised trial of elective continuous positive airway pressure (CPAP) verses rescue CPAP after extubation in infants following cardiac surgery

Study objectives

Elective use of nasal CPAP following tracheal extubation will reduce the incidence of the need for reintubation within 72 hours of primary tracheal extubation.

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

Neonatal Diseases

Interventions

Extubation of immediate nasal CPAP, extubation to face non-pressurised oxygen therapy only

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

13/02/2006

Completion date

12/02/2007

Eligibility

Key inclusion criteria

Not provided at time of registration

Participant type(s)

Patient

Age group

Neonate

Sex

Both

Target number of participants

110

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

13/02/2006

Date of final enrolment

12/02/2007

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Royal Brompton & Harefield NHS Trust

London

United Kingdom

SW3 6NP

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

Royal Brompton and Harefield NHS Trust (UK)

Funder Name

Clinical Research Committee

Funder Name

NHS R&D Support Funding

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