

CHEST - Pilot Study CPAP: Heliox Effects on Stability and Therapeutics

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| Submission date 19/07/2005 | Recruitment status Stopped | <input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol |
| Registration date 21/09/2005 | Overall study status Stopped | <input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results |
| Last Edited 04/01/2011 | Condition category Respiratory | <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 Parviz Habibi

Contact details
Imperial College London
Norfolk Place
London
United Kingdom
W21PG

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
AL001

Study information

Scientific Title

Acronym

CHEST

Study objectives

Heliox + oxygen driven continuous positive airway pressure (CPAP) reduces work of breathing and improves oxygenation and carbon dioxide clearance more than air + oxygen driven CPAP

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)

Treatment

Participant information sheet**Health condition(s) or problem(s) studied**

Respiratory distress requiring CPAP

Interventions

Heliox + oxygen driven continuous positive airway pressure (CPAP) versus air + oxygen driven CPAP.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Work of breathing

Secondary outcome measures

Oxygenation, carbon dioxide clearance

Overall study start date

01/09/2005

Completion date

01/03/2007

Reason abandoned (if study stopped)

Lack of funding

Eligibility

Key inclusion criteria

Children under 1 year of age with respiratory distress requiring CPAP

Participant type(s)

Patient

Age group

Child

Upper age limit

1 Years

Sex

Both

Target number of participants

20

Key exclusion criteria

Refusing consent, pneumothorax, $\text{FiO}_2 > 0.6$, imminent intubation, respiratory failure

Date of first enrolment

01/09/2005

Date of final enrolment

01/03/2007

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Imperial College London

London

United Kingdom
W21PG

Sponsor information

Organisation

Imperial College London (UK)

Sponsor details

Norfolk Place
London
England
United Kingdom
W21PG

Sponsor type

University/education

ROR

<https://ror.org/041kmwe10>

Funder(s)

Funder type

Industry

Funder Name

Gas will be provided by BOC Ltd. Equipment and application fees will be provided by Viasys

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