

# Randomised controlled trials and feasibility studies for newer modes of artificial ventilation in newborns with respiratory failure

<b>Submission date</b> 30/09/2005	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 30/09/2005	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
<b>Last Edited</b> 26/04/2011	<b>Condition category</b> 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

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### Contact details

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## Additional identifiers

### Protocol serial number

N0227097989

## Study information

Scientific Title

**Study objectives**

To assess the efficacy and safety of these newer modes of artificial ventilation, and assess their superiority over conventional ventilation in terms of mortality, analysis of cost of treatment, and short and long term respiratory outcome.

**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

**Study type(s)**

Treatment

**Health condition(s) or problem(s) studied**

Respiratory failure

**Interventions**

Randomised controlled trial of different modes of ventilation. Patients were randomised to different modes of ventilation, or the conventional ventilation methods.

**Intervention Type**

Other

**Phase**

Not Specified

**Primary outcome(s)**

Duration of ventilation and weaning

**Key secondary outcome(s))**

No secondary outcome measures

**Completion date**

31/07/2004

**Eligibility****Key inclusion criteria**

Babies less than 1500 g with respiratory distress syndrome (RDS) requiring ventilation

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Neonate

**Sex**

Not Specified

**Key exclusion criteria**

Not provided at time of registration

**Date of first enrolment**

01/01/1999

**Date of final enrolment**

31/07/2004

## **Locations**

**Countries of recruitment**

United Kingdom

England

**Study participating centre**

**Women & Children Division**

Middlesbrough

United Kingdom

TS4 3BW

## **Sponsor information**

**Organisation**

Department of Health

## **Funder(s)**

**Funder type**

Government

**Funder Name**

South Tees Hospitals 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?
<a href="#">Results article</a>	results	01/09/2006		Yes	No
<a href="#">Results article</a>	results	01/03/2007		Yes	No
<a href="#">Results article</a>	results	01/03/2009		Yes	No
<a href="#">Results article</a>	results	01/09/2009		Yes	No