

A double-blind randomised controlled trial of nasal continuous positive airway pressure (CPAP) in patients with obstructive sleep apnoea

Submission date 23/01/2004	Recruitment status No longer recruiting	☐ Prospectively registered
Registration date 23/01/2004	Overall study status Completed	☐ Protocol
Last Edited 21/09/2011	Condition category Respiratory	☐ Statistical analysis plan
		☐ Results
		☐ Individual participant data
		☐ Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Additional identifiers

Protocol serial number

R/32/01.98/Kendrick/S

Study information

Scientific Title

A double-blind randomised controlled trial of nasal continuous positive airway pressure (CPAP) in patients with obstructive sleep apnoea

Study objectives

Does nasal Continuous Positive Airways Pressure (CPAP) significantly reduce symptoms in patients with obstructive sleep apnoea?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Double blind randomised controlled trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Respiratory tract diseases: Obstructive sleep apnoea

Interventions

Not provided at time of registration

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

1. Number of hours of CPAP used per night
2. The mean number of hours usage per week
3. Epworth Score

Key secondary outcome(s))

1. SF-36
2. Steer Clear driving simulation test
3. Sleep-wake diary data

Completion date

31/12/2001

Eligibility

Key inclusion criteria

Not provided at time of registration

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

05/10/1998

Date of final enrolment

31/12/2001

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

United Bristol Health Care Trust

Bristol

United Kingdom

BS1 6SY

Sponsor information**Organisation**

NHS R&D Regional Programme Register - Department of Health (UK)

Funder(s)

Funder type

Government

Funder Name

NHS Executive South West (UK)

Results and Publications

Individual participant data (IPD) sharing plan**IPD sharing plan summary**

Not provided at time of registration