Effect of mouth taping at night on asthma control

Submission date Recruitment status Prospectively registered 28/09/2005 No longer recruiting [] Protocol [] Statistical analysis plan Registration date Overall study status 15/11/2005 Completed [X] Results Individual participant data **Last Edited** Condition category 15/07/2010 Respiratory

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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers TC03/5

Study information

Scientific Title

Study objectives

Taping the mouth at night to ensure nose breathing will improve asthma control

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)

Other

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Asthma

Interventions

Crossover design of taping mouth at night versus usual breathing

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

Morning peak expiratory flow and symptom scores

Secondary outcome measures

- 1. Evening peak flow
- 2. Diurnal variation in peak flow
- 3. FEV1
- 4. Beta-agonist use
- 5. Night time waking

- 6. Quality of life
- 7. Acceptability
- 8. Compliance with taping

Overall study start date

27/05/2004

Completion date

31/05/2006

Eligibility

Key inclusion criteria

- 1. Asthma
- 2. Age 18-72
- 3. Forced expiratory volume in one second (FEV1) over 50% predicted
- 4. Nocturnal or early morning symptoms
- 5. Pro re nata (prn) (as needed) beta-agonist use

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Upper age limit

72 Years

Sex

Both

Target number of participants

60

Key exclusion criteria

- 1. Previous training in Buteyko technique
- 2. Inability to breathe through nose
- 3. Known sleep apnoea
- 4. Over 10 pack years smoking history
- 5. Other significant uncontrolled disease

Date of first enrolment

27/05/2004

Date of final enrolment

31/05/2006

Locations

Countries of recruitment

England

United Kingdom

Study participating centre
Division Respiratory Medicine
Nottingham
United Kingdom
NG5 1PB

Sponsor information

Organisation

University of Nottingham (UK)

Sponsor details

Research Innovation Services
University Park
Nottingham
England
United Kingdom
NG7 2RD
paul.cartledge@nottingham.ac.uk

Sponsor type

University/education

ROR

https://ror.org/01ee9ar58

Funder(s)

Funder type

Charity

Funder Name

British Lung Foundation (UK) -)ref: TC03/5)

Alternative Name(s)

Funding Body Type

Private sector organisation

Funding Body Subtype

Trusts, charities, foundations (both public and private)

Location

United Kingdom

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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/06/2009		Yes	No