

Randomised controlled trial to compare 2 different methods of nasendoscopic examination of the hypopharynx

Submission date 29/09/2006	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 29/09/2006	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 30/08/2013	Condition category Ear, Nose and Throat	<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

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0301167003

Study information

Scientific Title

Study objectives

To compare 2 different modified valsalva techniques as aids to nasendoscopic examination of the hypopharynx.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

CT

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Other

Participant information sheet

Health condition(s) or problem(s) studied

Ear, Nose and Throat: Throat disorder

Interventions

Methods of nasendoscopic examination of the hypopharynx 1 vs method 2

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Percentage visualisation of 4 subsites: post-cricoid, right and left pyriform fossae & upper oesophagus. Patient discomfort.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/07/2005

Completion date

01/10/2005

Eligibility

Key inclusion criteria

50-100 outpatients >18 years attending the Otolaryngology clinic suffering a throat disorder that would require an examination of the hypopharynx with a nasendoscope.

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Not Specified

Target number of participants

100

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/07/2005

Date of final enrolment

01/10/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Department of Otolaryngology
Blackpool

United Kingdom
FY3 8NR

Sponsor information

Organisation

Record Provided by the NHSTCT Register - 2006 Update - Department of Health

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

Blackpool Fylde & Wyre Hospitals NHS Trust (UK)

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

