ISRCTN40986271 https://doi.org/10.1186/ISRCTN40986271

The effect of timing of the application of topical anaesthetic prior to rigid nasendoscopy

Submission date	Recruitment status	[_] Prospectively registered
29/09/2006	No longer recruiting	[_] Protocol
Registration date	**Overall study status**	[_] Statistical analysis plan
29/09/2006	Completed	[X] Results
Last Edited	**Condition category**	[_] Individual participant data
08/12/2010	Surgery	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0212177261

Study information

Scientific Title

Study objectives
Is it best to wait one or ten minutes before performing nasendoscopy after the application of cophenylcaine?

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)
Not Specified

Participant information sheet

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

Interventions
Quantitative, randomised controlled trial
Treatment 1: wait 1 minute
Treatment 2: wait 10 minutes

Intervention Type
Procedure/Surgery

Phase
Not Specified

Primary outcome measure
Visual analogue scales of pain and discomfort for patients.

Secondary outcome measures

Not provided at time of registration

Overall study start date
01/03/2006

Completion date
01/09/2006

Eligibility

Key inclusion criteria
Patients who, by the decision of the examining clinician, need to undergo examination of their nasal cavity using rigid nasendoscopy.

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
50

Key exclusion criteria
1. Unable to provide informed consent
2. Patients in whom lignocaine or phenylphrine is contra indicated for medical reasons
3. Patients in whom only one nasal cavity or part of one cavity is to be examined

Date of first enrolment
01/03/2006

Date of final enrolment
01/09/2006

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

ENT
Bath
United Kingdom
BA1 3NG

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
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Sponsor type
Government

Website
http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type
Government

Funder Name
Royal United Hospital Bath NHS Trust

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	01/03/2007		Yes	No