ISRCTN86425765 https://doi.org/10.1186/ISRCTN86425765

Is there any advantage using water as a lubricant when performing flexible 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
06/04/2009	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
N0106163114

Study information

Scientific Title

Study objectives
To establish what type of lubrication is most beneficial when performing nasendoscopy.

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Single blind prospective randomised controlled trial

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
Not specified

Study type(s)
Treatment

Participant information sheet

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

Interventions
Group A: water as lubricant
Group B: petroleum jelly as lubricant

Intervention Type
Procedure/Surgery

Phase
Not Specified

Primary outcome measure
1. Visual analogue score (VAS) of patient discomfort
2. VAS of picture quality information gained by the doctor
3. VAS of the ease of passage of the nasendoscope

Secondary outcome measures

Not provided at time of registration

Overall study start date
01/06/2005

Completion date
01/12/2005

Eligibility

Key inclusion criteria
Patients requiring nasendoscopy

Participant type(s)
Patient

Age group
Adult

Sex
Not Specified

Target number of participants
150 - 75 in each group.

Key exclusion criteria
1. Patients unable to consent
2. Allergic to lubricant
3. Unable to complete VAS
4. Patients who require local anaesthetic

Date of first enrolment
01/06/2005

Date of final enrolment
01/12/2005

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

ENT Department
Gloucester
United Kingdom
GL1 3NN

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
Gloucestershire R&D Consortium (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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/04/2006		Yes	No