ISRCTN24006995 https://doi.org/10.1186/ISRCTN24006995

Nasal decongestant study: a prospective double-blind randomised comparative study of nasal decongestant and vasoconstrictive efficacy of two commonly used preparations, namely cocaine and adrenaline, with normal saline and adrenaline with normal saline

Submission date
28/09/2007

Recruitment status
No longer recruiting

Registration date
28/09/2007

Overall study status
Completed

Last Edited
05/06/2014

Condition category
Surgery

[ ] Prospectively registered
[ ] Protocol
[ ] 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

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0247184156

Study information

Scientific Title

Study objectives
During endoscopic nasal surgery a pre-operative topical decongestion of the nose is achieved in the theatre by instillation of a decongestant solution. During operation this solution is used to achieve haemostasis. This research aims to compare the efficacy of two preparations and justify the use of cocaine.

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)
Hospital

Study type(s)
Treatment

Participant information sheet

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

Interventions
Cocaine and adrenaline with normal saline and adrenaline with normal saline

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)
Cocaine, adrenaline, normal saline

Primary outcome measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/08/2006

Completion date
01/11/2006

Eligibility

Key inclusion criteria
Not provided at time of registration

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
Not provided at time of registration

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/08/2006

Date of final enrolment
01/11/2006

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Swindon & Marlborough NHS Trust
Swindon
United Kingdom
SN3 6BB

Sponsor information

Organisation
Record Provided by the NHSTCT Register - 2007 Update - Department of Health

Sponsor details
The Department of Health, Richmond House, 79 Whitehall
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dhmail@doh.gsi.org.uk

Sponsor type
Government

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

Funder(s)

Funder type
Government

Funder Name
Swindon and Marlborough 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