

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	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 28/09/2007	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 05/06/2014	Condition category Surgery	<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

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
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

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