

A randomised controlled trial of local anaesthetic in merocel nasal pack removal

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 07/09/2015	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

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0015163350

Study information

Scientific Title

A randomised controlled trial of local anaesthetic in merocel nasal pack removal

Study objectives

Is soaking merocel nasal packs in local anaesthetic with adrenaline more effective in reducing pain and bleeding than soaking the pads in saline?

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

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Ear, Nose and Throat: Nasal packing

Interventions

Local anaesthetic with adrenaline vs saline

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

adrenaline

Primary outcome measure

Pain score upon removal of packs and again at timed intervals after removal.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/03/2005

Completion date

01/12/2005

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

50

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/03/2005

Date of final enrolment

01/12/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

North Cheshire Hospitals NHS Trust
Warrington
United Kingdom
WA5 1QG

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
+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

North Cheshire Hospitals NHS Trust (UK), NHS R&D Support Funding

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