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A Pilot comparative study of the efficacy and patient acceptability of the Rapid Rhino tamponade balloon and Merocel nasal tampon in acute epistaxis

Submission date
30/09/2004

Recruitment status
No longer recruiting

Registration date
30/09/2004

Overall study status
Completed

Last Edited
20/08/2015

Condition category
Signs and Symptoms

[ ] 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
N0083126301

Study information

Scientific Title
A Pilot comparative study of the efficacy and patient acceptability of the Rapid Rhino tamponade balloon and Merocel nasal tampon in acute epistaxis

Study objectives
There is no difference between the rapid rhino tamponade balloon and the merocel nasal tampon in preventing bleeding from the nose.

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)
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
Signs and Symptoms: Acute epistaxis

Interventions
Rapid rhino tamponade balloon vs merocel nasal tampon

Intervention Type
Device

Primary outcome measure

1. Time to cessation of bleeding
2. Patient comfort

Secondary outcome measures
Not provided at time of registration

Overall study start date
13/03/2003

Completion date
12/09/2003

Eligibility

Key inclusion criteria
126 patients

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
126

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
13/03/2003

Date of final enrolment
12/09/2003

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Calderdale and Huddersfield NHS Trust
Halifax

United Kingdom
HX3 0PW

Sponsor information

Organisation
Department of Health

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

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

Funder(s)

Funder type
Government

Funder Name
Calderdale and Huddersfield 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