

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

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| Submission date 30/09/2004 | Recruitment status No longer recruiting | <input type="checkbox"/> Prospectively registered |
| | | <input type="checkbox"/> Protocol |
| Registration date 30/09/2004 | Overall study status Completed | <input type="checkbox"/> Statistical analysis plan |
| | | <input type="checkbox"/> Results |
| Last Edited 20/08/2015 | Condition category Signs and Symptoms | <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

Protocol serial number

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

Study type(s)

Treatment

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(s)

1. Time to cessation of bleeding
2. Patient comfort

Key secondary outcome(s)

Not provided at time of registration

Completion date

12/09/2003

Eligibility

Key inclusion criteria

126 patients

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

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**

United Kingdom

England

Study participating centre

Calderdale and Huddersfield NHS Trust

Halifax

United Kingdom

HX3 0PW

Sponsor information**Organisation**

Department of Health

Funder(s)**Funder type**

Government

Funder Name

Calderdale and Huddersfield NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration