

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

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

### Contact name

Mr Dominic Martin-Hirsch

### Contact details

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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

**Study outputs**

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Participant information sheet</a>	Participant information sheet	11/11/2025	11/11/2025	No	Yes