

# The use of Sambucus Comp. in the treatment of postnasal drip

<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> 31/03/2020	<b>Condition category</b> 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

**Protocol serial number**  
N0176127651

## Study information

**Scientific Title**  
The use of Sambucus Comp. in the treatment of postnasal drip

**Study objectives**

The aim of the study is to ascertain whether the anthroposophic medicine, Sambucus Comp. has a role in the treatment of postnasal drip.

**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**

Ear, Nose and Throat

**Interventions**

Sambucus Comp. vs standard practice

**Intervention Type**

Drug

**Phase**

Not Applicable

**Drug/device/biological/vaccine name(s)**

Sambucus Comp.

**Primary outcome(s)**

Not provided at time of registration

**Key secondary outcome(s)**

Not provided at time of registration

**Completion date**

01/04/2004

**Eligibility****Key inclusion criteria**

Not provided at time of registration

**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**

01/03/2003

**Date of final enrolment**

01/04/2004

**Locations****Countries of recruitment**

United Kingdom

England

**Study participating centre**

Radcliffe Infirmary

Oxford

United Kingdom

OX2 6HE

**Sponsor information****Organisation**

Department of Health

**Funder(s)****Funder type**

Hospital/treatment centre

**Funder Name**

Oxford Radcliffe Hospitals NHS Trust (UK)

**Results and Publications**

Individual participant data (IPD) sharing plan

**IPD sharing plan summary**

Not provided at time of registration