

# Microsuction versus instruments for dewaxing: a randomised controlled trial

<b>Submission date</b> 28/09/2007	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
<b>Registration date</b> 28/09/2007	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
<b>Last Edited</b> 28/03/2012	<b>Condition category</b> Ear, Nose and Throat	<input type="checkbox"/> Individual participant data

**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**  
N0212190355

## Study information

**Scientific Title**

**Study objectives**

There are two methods of wax removal in ENT clinic. The use of microsuction and the use of instruments. We aim to see if either method causes less discomfort and or complications.

**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: Ear dewaxing

**Interventions**

Quantitative randomised controlled trial. Questionnaire based study. A total of 100 patients are required.

**Intervention Type**

Other

**Phase**

Not Specified

**Primary outcome(s)**

Not provided at time of registration

**Key secondary outcome(s)**

Not provided at time of registration

**Completion date**

01/07/2008

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

1. Any patient requiring wax removal from their ear in ENT clinic
2. Without otitis externa, mastoid cavity

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Not Specified

**Sex**

Not Specified

**Key exclusion criteria**

Unable/unwilling to consent.

**Date of first enrolment**

01/07/2006

**Date of final enrolment**

01/07/2008

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

United Kingdom

England

**Study participating centre**

ENT

Bath

United Kingdom

BA1 3NG

**Sponsor information****Organisation**

Record Provided by the NHSTCT Register - 2007 Update - Department of Health

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

Government

**Funder Name**

## 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="#">Results article</a>	results	01/10/2006		Yes	No