

# A randomised controlled trial of outpatient hysteroscopy looking at the effects of temperature and pressure

<b>Submission date</b> 30/09/2005	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
<b>Registration date</b> 30/09/2005	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
<b>Last Edited</b> 18/10/2017	<b>Condition category</b> Surgery	<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

Dr Adam L Magos

### Contact details

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## Additional identifiers

### Protocol serial number

N0256159467

## Study information

### Scientific Title

A randomised controlled trial of outpatient hysteroscopy looking at the effects of temperature and pressure

### **Study objectives**

To determine whether women who undergo hysteroscopy experience more discomfort when there is a change in pressure and temperature of the saline used to distend the endometrial cavity

### **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)**

Other

### **Health condition(s) or problem(s) studied**

Surgery: Hysteroscopy

### **Interventions**

Randomised controlled trial:

1. Normal saline, 37degree celsius, 150 mmHg
2. Normal saline, room temperature 100mmHg
3. Normal saline, room temperature, 200 mmHg
4. Saline, room temperature, 150 mmHg

### **Intervention Type**

Procedure/Surgery

### **Phase**

Not Specified

### **Primary outcome(s)**

Service outcome development

### **Key secondary outcome(s)**

Not provided at time of registration

### **Completion date**

15/07/2004

## **Eligibility**

**Key inclusion criteria**

120 patients

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Adult

**Sex**

Female

**Key exclusion criteria**

Does not match inclusion criteria

**Date of first enrolment**

15/07/2003

**Date of final enrolment**

15/07/2004

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

United Kingdom

England

**Study participating centre**

University Department Of Obstetrics and Gynaecology

London

United Kingdom

NW3 2QG

**Sponsor information****Organisation**

Department of Health

**Funder(s)**

**Funder type**

Hospital/treatment centre

**Funder Name**

The Royal Free Hampstead NHS Trust (UK) NHS R&D Support Funding

## **Results and Publications**

**Individual participant data (IPD) sharing plan****IPD sharing plan summary**

Not provided at time of registration