

# A prospective randomised comparison of surgical treatment and expectant management of intrauterine polyps/fibroids in premenopausal women with abnormal uterine bleeding in an office hysteroscopy setting

<b>Submission date</b> 30/09/2004	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
<b>Registration date</b> 30/09/2004	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 19/08/2015	<b>Condition category</b> Urological and Genital Diseases	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<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 Kevin Phillips

**Contact details**  
Department of Women's Health  
Castle Hill Hospital  
Castle Road  
Cottingham  
United Kingdom  
HU16 5JQ  
+44 (0)1842 875875  
abc@email.com

## Additional identifiers

**Protocol serial number**  
N0084125088

# Study information

## Scientific Title

A prospective randomised comparison of surgical treatment and expectant management of intrauterine polyps/fibroids in premenopausal women with abnormal uterine bleeding in an office hysteroscopy setting

## Study objectives

To determine the effect of treatment on the symptoms experienced and to determine the natural history of polyps/fibroids in relation to the symptoms in patients who have intrauterine lesion in situ.

## 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

Urological and Genital Diseases: Polyps/fibroids

## Interventions

Patients will be randomised preoperatively by a computer generated sequence into one of the two study arms: Hysteroscopic treatment or diagnostic hysteroscopy alone.

The control group will comprise of asymptomatic patients undergoing sterilisation who will have diagnostic hysteroscopy.

## Intervention Type

Procedure/Surgery

## Primary outcome(s)

Not provided at time of registration

## Key secondary outcome(s)

Not provided at time of registration

## Completion date

01/12/2005

## Eligibility

**Key inclusion criteria**

Women attending the clinic for routine check up.

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Adult

**Sex**

Female

**Key exclusion criteria**

Not provided at time of registration

**Date of first enrolment**

08/04/2003

**Date of final enrolment**

01/12/2005

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

United Kingdom

England

**Study participating centre**

Castle Hill Hospital

Cottingham

United Kingdom

HU16 5JQ

**Sponsor information****Organisation**

Department of Health

**Funder(s)**

**Funder type**

Government

**Funder Name**

The North and South Bank Research and Development Consortium (UK)

## **Results and Publications**

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

Not provided at time of registration