

# Does the use of local anaesthetic make trans-rectal ultrasound guided biopsy more comfortable. To evaluate the pain experienced by the patient when having this procedure

<b>Submission date</b> 30/09/2005	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
<b>Registration date</b> 30/09/2005	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 27/04/2018	<b>Condition category</b> Surgery	<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

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## Additional identifiers

**EudraCT/CTIS number**

**IRAS number**

**ClinicalTrials.gov number**

**Secondary identifying numbers**

N0436146667

## **Study information**

### **Scientific Title**

Does the use of local anaesthetic make trans-rectal ultrasound guided biopsy more comfortable.  
To evaluate the pain experienced by the patient when having this procedure

### **Study objectives**

To assess whether the use of local anaesthetic makes trans-rectal ultrasound guided biopsy more comfortable. To evaluate the pain experienced by the patient when having this procedure.

### **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

### **Secondary study design**

Randomised controlled trial

### **Study setting(s)**

Hospital

### **Study type(s)**

Diagnostic

### **Participant information sheet**

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

Surgery: Anaesthesia

### **Interventions**

Randomised controlled trial

### **Intervention Type**

Procedure/Surgery

### **Phase**

Not Specified

**Primary outcome measure**

Not provided at time of registration

**Secondary outcome measures**

Not provided at time of registration

**Overall study start date**

01/08/2002

**Completion date**

01/06/2003

## Eligibility

**Key inclusion criteria**

Male patients referred for trans-rectal ultrasound guided biopsy as an outpatient to St James's

**Participant type(s)**

Patient

**Age group**

Adult

**Sex**

Male

**Target number of participants**

50

**Key exclusion criteria**

1. Patients who are unable to answer the questionnaire (at the discretion of the radiographer, eg dementia, mental illness, language difficulties)
2. Patients not suitable for trans-rectal ultrasound biopsy

**Date of first enrolment**

01/08/2002

**Date of final enrolment**

01/06/2003

## Locations

**Countries of recruitment**

England

United Kingdom

**Study participating centre**

**St James's University Hospital, Radiology**  
Lincoln Wing  
Beckett Street  
Leeds  
United Kingdom  
LS9 7TF

## **Sponsor information**

### **Organisation**

Department of Health

### **Sponsor details**

Richmond House  
79 Whitehall  
London  
United Kingdom  
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+44 (0)20 7307 2622  
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### **Sponsor type**

Government

### **Website**

<http://www.dh.gov.uk/Home/fs/en>

## **Funder(s)**

### **Funder type**

Government

### **Funder Name**

Leeds Teaching Hospitals NHS Trust (UK)

## **Results and Publications**

### **Publication and dissemination plan**

Not provided at time of registration

### **Intention to publish date**

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

### **IPD sharing plan summary**

Not provided at time of registration