ISRCTN77665682 https://doi.org/10.1186/ISRCTN77665682

Combining surface and infiltration anaesthesia for Transrectal Ultrasonography of the Prostate (TRUS) biopsy: is it better?

Submission date 30/09/2005	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [_] Results [_] Individual participant data [_] Record updated in last year
Registration date 30/09/2005	**Overall study status** Completed	
Last Edited 11/04/2014	**Condition category** Surgery	

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
N0155153512

Study information

Scientific Title

Study objectives
Is combining surface and infiltration anaesthesia for Transrectal Ultrasonography of the Prostate (TRUS) biopsy more effective in relieving the pain of the procedure?

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Double blind randomised controlled trial

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
Not specified

Study type(s)
Not Specified

Participant information sheet

Health condition(s) or problem(s) studied
Surgery: Anaesthesia

Interventions
The intervention is infiltration of local anaesthesia in periprostatic tissue with 1% lignocaine solution: and the ano-rectal mucosa anaesthetized by application of 2% lignocaine jelly.

Intervention Type
Procedure/Surgery

Phase
Not Specified

Primary outcome measure
Pain relief by visual analogue score

Secondary outcome measures
Not provided at time of registration

Overall study start date
07/07/2004

Completion date
30/09/2004

Eligibility

Key inclusion criteria
80 patients undergoing Transrectal Ultrasonography of the Prostate (TRUS) biopsy

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
80

Key exclusion criteria
Known allergy to lignocaine.

Date of first enrolment
07/07/2004

Date of final enrolment
30/09/2004

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Urology Unit
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Sponsor information

Organisation
Department of Health

Sponsor details
Richmond House
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Sponsor type
Government

Website
http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type
Government

Funder Name
Pennine Acute Hospitals NHS Trust (UK), NHS R&D Support Funding Own Account

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